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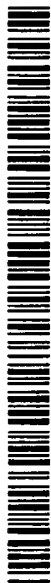


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- (71) Applicant: GUILFORD PHARMACEUTICALS [US/US]; 6611 Tributary Street, Baltimore, MD 21224 (US).
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- (72) Inventors: JACKSON, Paul, F.; 102 Glenmore Court, Bel Air, MD 21014 (US). SLUSHER, Barbara, S.; 7424 Longfield Drive, Kingsville, MD 21087 (US).
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.
- (74) Agent: SUET, M., Chong; Lyon & Lyon LLP, 633 West Fifth Street, Suite 4700, Los Angeles, CA 90071-2066 (US).



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(54) Title: METABOTROPIC GLUTAMATE RECEPTOR LIGAND DERIVATIVES AS NAALADase INHIBITORS

(57) Abstract: The present invention relates to metabotropic glutamate receptor ligand derivatives and methods of using the same to inhibit NAALADase enzyme activity, to effect neuronal activities, to inhibit angiogenesis, and to treat glutamate abnormalities, compulsive disorders, pain, diabetic neuropathy, and prostate diseases, as well as pharmaceutical compositions comprising the same.

**METABOTROPIC GLUTAMATE RECEPTOR LIGAND DERIVATIVES**  
**AS NAALADase INHIBITORS**

**BACKGROUND OF THE INVENTION**

5           The present invention relates to metabotropic glutamate receptor ligand derivatives, pharmaceutical compositions comprising such derivatives and methods of their use to inhibit NAALADase enzyme activity, thereby effecting neuronal activities, inhibiting angiogenesis and  
10   treating glutamate abnormalities, compulsive disorders, pain, diabetic neuropathy, and prostate diseases. When a metabotropic glutamate receptor ligand, preferably an mGluR3 receptor ligand, is attached to a metal chelating group capable of interacting with the metal atom(s) at the  
15   active site of NAALADase, it is expected that the resulting compound will be a potent and specific NAALADase inhibitor.

          The NAALADase enzyme, also known as prostate specific membrane antigen (PSM or PSMA) and human glutamate carboxypeptidase II (GCP II), catalyzes the hydrolysis of  
20   the neuropeptide N-acetyl-aspartyl-glutamate ("NAAG") to N-acetyl-aspartate ("NAA") and glutamate. Based upon amino acid sequence homology, NAALADase has been assigned to the M28 family of peptidases. There is, as yet, no crystallographic evidence of the structure of the NAALADase  
25   enzyme.

          Recent studies have implicated NAALADase in the pathogenesis of glutamate-mediated disorders. Neuropathological studies on post-mortem tissue from patients with amyotrophic lateral sclerosis (ALS) indicate  
30   large decreases of N-acetylaspartate (NAA) and N-acetylaspartylglutamate (NAAG) tissue concentrations occurring in association with neuronal degeneration, and increases of NAA and NAAG in cerebral spinal fluid (CSF) from patients with ALS. Concordantly, abnormal NAAG levels  
35   and NAALADase activity have also been observed in post-mortem prefrontal and limbic brain tissue of schizophrenic

patients. Autopsy studies also suggest a strong correlation between NAAG/NAA and Alzheimer's disease. In post-mortem brain tissue, NAA and NAAG levels were found to be selectively decreased in brain areas (hippocampus and amygdala) affected by Alzheimer's disease pathology.

Glutamate serves as the predominant excitatory neurotransmitter in the central nervous system (CNS). Neurons release glutamate in greater quantities when they are deprived of oxygen, as may occur during an ischemic brain insult such as a stroke or a heart attack. This excess release of glutamate in turn causes over-stimulation (excitotoxicity) of N-methyl-D-aspartate (NMDA), AMPA, Kainate and MGR receptors. When glutamate binds to these receptors, ion channels in the receptors open, permitting flows of ions across their cell membranes, e.g.,  $\text{Ca}^{2+}$  and  $\text{Na}^+$  into the cells and  $\text{K}^+$  out of the cells. These flows of ions, especially the influx of  $\text{Ca}^{2+}$ , cause over-stimulation of the neurons. The over-stimulated neurons secrete more glutamate, creating a feedback amplification effect which is believed to ultimately result in cell death via the production of proteases, lipases, and free radicals.

Excessive activation of glutamate receptors has been implicated in various neurological diseases and conditions, including spinal cord injury, epilepsy, stroke, Alzheimer's disease, Parkinson's Disease, Amyotrophic Lateral Sclerosis (ALS), Huntington's Disease, schizophrenia, acute and chronic pain, ischemia and neuronal loss following hypoxia, hypoglycemia, ischemia, trauma, nervous insult, compulsive disorders (particularly drug and alcohol dependence), demyelinating diseases, peripheral neuropathies, and diabetic neuropathy.

In particular, glutamatergic abnormalities have been associated with schizophrenia. For example, phencyclidine (PCP) and other antagonists of N-methyl-D-aspartate (NMDA) receptors induce psychotomimetic properties in healthy

individuals and exacerbate preexisting symptoms of schizophrenia, suggesting that a depression of glutamate transmission might contribute to schizophrenia. Additionally, it has been reported that antagonists of non-NMDA receptors or pretreatments that attenuate glutamate release reduce mnemonic and other behavioral effects of NMDA receptor antagonists. Studies have also shown that stimulation of certain subtypes of mGlu receptors mediates presynaptic depression and decreases evoke release of glutamate.

Recent studies have also advanced a glutamatergic basis for compulsive disorders, particularly drug dependence. For example, neurophysiological and pathological effects of ethanol have been found to be mediated through the glutamatergic system. Specifically, acute exposure to ethanol disrupts glutamatergic neurotransmission by inhibiting ion flow through channels in glutamate receptors, whereas chronic exposure up-regulates the number of glutamate receptors and thereby increases ion flow. Acute withdrawal from ethanol results in hyperexcitability and seizures in the presence of up-regulated channels, thereby making postsynaptic neurons vulnerable to excitotoxic damage.

Post mortem examinations of histologically normal brains from alcoholics have shown that chronic alcoholism moderately increases the density of the NMDA subtype of glutamate receptors in the frontal cortex. This up-regulation may represent a stage of ethanol-induced chronic neurotoxicity. As such, neurobiological effects of alcoholism, including intoxication, withdrawal seizures, delirium tremens, Wernicke-Korsakoff syndrome and fetal alcohol syndrome, can be understood as a spectrum of the consequences of ethanol's effect on the glutamatergic system. In this regard, alcoholism may be considered another member of the expanding family of glutamate-related

neurological disorders.

The glutamatergic system has also been implicated in the behavioral effects of other abused drugs. For example, studies have shown that glutamatergic antagonists block  
5 motor-stimulating activities induced by amphetamine and cocaine, and glutamatergic agonists cause the same stereotypy as that produced by amphetamine. These results represent pharmacological evidence that the expression of the stereotypic effect of psychomotor stimulants involves  
10 the glutamatergic system.

Epidemiologic studies have revealed a strong correlation between drug dependence and other compulsive disorders. Additionally, a common genetic anomaly has been found among people with alcoholism, cocaine dependence,  
15 nicotine dependence, pathological gambling, attention deficit disorder (ADD), Tourette's syndrome, compulsive overeating and obesity. Such disorders are believed to be manifestations of the effects of excitotoxicity.

Based on the above findings, the present inventors  
20 tested and found NAALADase inhibitors to be efficacious in the pharmacotherapy of glutamate abnormalities, such as drug dependence, diabetic neuropathy, pain, schizophrenia, ischemic injury, and anxiety.

Ischemic injury may occur as a focal or global  
25 disruption of blood supply. Following ischemic insult, widespread neuronal depolarization occurs. Depolarization stimulates release of the stored neurotransmitter glutamate and results in impaired capacity of glutamate uptake mechanisms. Impaired glutamate uptake and enhanced  
30 glutamate release contribute to sustained elevation of extracellular glutamate in ischemic tissue, and may result in tissue damage. As more damage occurs, more glutamate may be released. Although not limited to any particular theory, it is believed that by interfering with or  
35 eliminating this cascade of glutamate toxicity, the

compositions and methods of the present convention may be clinically useful in curbing the progression of ischemic injury.

5       Afferent pain fibers of the A- $\delta$  and C types have their primary cell bodies in the dorsal root ganglia; central extensions of these nerve cells project, via the dorsal root, to the dorsal horn of the spinal cord or to the nucleus of the trigeminal nerve; the peripheral terminations of these primary pain receptors are the branch  
10       nerve endings in the skin and other organs. Excitatory amino acids, including glutamate, and ATP are putative neurotransmitters at the dorsal horn terminus of primary A- $\delta$  fibers. The conscious awareness or perception of pain occurs only when the pain impulses actually reach the  
15       thalamocortical level. Although not limited to any particular theory, it is believed that by interfering with such nerve impulses, the compositions and methods of the present convention may be clinically useful in limiting or eliminating pain.

20       Diabetic neuropathy is a slowly progressive, mixed sensorimotor and autonomic neuropathy. A variety of pathogenic mechanisms have been proposed for diabetic neuropathy, including alteration in nerve metabolism induced by ischemia and, in some cases, autoimmunity.  
25       Although not limited to any particular theory, it is believed that by interfering with or eliminating these effects, the compositions and methods of the present convention may be clinically useful in curbing the progression of diabetic neuropathy.

30       Excessive activation of glutamate receptors has been implicated in anxiety and anxiety disorders. Significantly higher glutamate plasma levels have been detected in patients with mood disorders than in comparison subjects. Studies also suggest that the pharmacological effect of  
35       anxiolytic agents is mediated through the glutamatergic

system. Although not limited to any particular theory, it is believed that by interfering with or eliminating these effects, the compositions and methods of the present convention may be clinically useful in curbing anxiolytic activity.

Most research and development activity to date have focused on blocking post-synaptic glutamate receptors with compounds such as NMDA antagonists, glycine antagonists, and other post-synaptic excitatory amino acid (EAA) receptor blockers. Unfortunately, these efforts have proven difficult because each receptor has multiple sites to which glutamate may bind; in addition, these agents produce severe toxicities even under normal conditions, thus limiting their clinical use. Although not limited to any one particular theory, it is believed that NAALADase inhibitors block glutamate release pre-synaptically without interacting with post-synaptic glutamate receptors. Since NAALADase inhibitors do not appear to alter basal glutamate levels, they may be devoid of the behavioral toxicities associated with post-synaptic glutamate antagonists.

In addition to glutamate, NAALADase has also been associated with prostate-specific membrane antigen (PSMA). In particular, it has been shown that PSMA cDNA confers NAALADase activity and that NAALADase and PSMA exhibit at least 86% homologous sequence identity. Carter et al., *Proc. Natl. Acad. Sci.*, Vol. 93, pp. 749-753 (1996). The molecular cloning of PSMA has been reported as a potential prostate carcinoma marker and hypothesized to serve as a target for imaging and cytotoxic treatment modalities for prostate cancer. Additionally, PSMA antibodies, particularly indium-111 labelled and itrium labelled PSMA antibodies, have been described and examined clinically for the diagnosis and treatment of prostate cancer. PSMA is expressed in prostatic ductal epithelium and is present in seminal plasma, prostatic fluid and urine.

Applicants have found NAALADase inhibitors to be effective in treating prostate diseases, particularly prostate cancer. Although not limited to any particular theory, it is believed that NAALADase inhibitors inhibit PSMA activity. Since mAbs to PSMA have been found to target 23 non-prostate carcinomas (Lui et al., *Science Research*, Vol. 57, pp. 3629-34 (1997)), the present inventors hypothesize that NAALADase inhibitors would also be effective in treating non-prostate cancers, particularly in tissues where NAALADase resides, such as the brain, kidney and testis.

NAALADase has also been found in neovasculature (new blood vessels). The present inventors have discovered that NAALADase inhibitors inhibit or prevent growth of neovasculature (angiogenesis), thereby providing potential therapeutic applications in treating diseases dependent upon angiogenesis. Examples of angiogenesis-dependent diseases include without limitation rheumatoid arthritis, cardiovascular disease, neovascular diseases of the eye, peripheral vascular disorders, and dermatologic ulcers. Angiogenesis is also essential for normal physiological processes, such as growth, fertility and soft tissue wound healing.

Cancer is another disease dependent upon angiogenesis. Cancer tumor cells secrete or release angiogenic substances that activate nearby endothelial cells. These endothelial cells respond by expressing a cell autonomous pattern of behavior that culminates in the formation of new blood vessels. Since research has demonstrated that angiogenesis is necessary to sustain the growth, invasion and metastasis of cancer tumors, the neovasculature inhibiting activity of NAALADase inhibitors further supports their utility in treating all types of cancers.

While a few NAALADase inhibitors have been identified, they have only been used in non-clinical research.

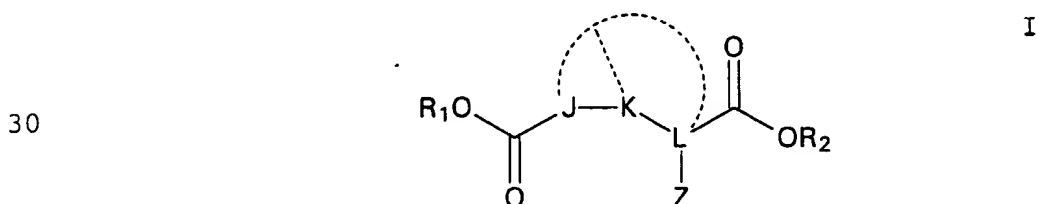


Examples of such inhibitors include general metallopeptidase inhibitors such as o-phenanthroline, metal chelators such as EGTA and EDTA, and peptide analogs such as quisqualic acid and  $\beta$ -NAAG. These compounds either have  
5 toxic side effects or are incapable of being administered in pharmaceutically effective amounts. NAAG is an agonist at group II metabotropic glutamate receptors, specifically mGluR3 receptors. When a metabotropic glutamate receptor ligand, preferably an mGluR3 receptor ligand, is attached  
10 to a metal chelating group capable of interacting with the metal atom(s) at the active site of NAALADase, it is expected that the resulting compound will be a potent and specific NAALADase inhibitor. In view of the broad range of potential applications, a need exists for new NAALADase  
15 inhibitors, pharmaceutical compositions comprising such inhibitors, and methods of their use.

#### SUMMARY OF THE INVENTION

The present invention relates to metabotropic glutamate receptor ligand compounds and compositions  
20 useful for inhibiting N-Acetylated  $\alpha$ -Linked Acidic Dipeptidase (NAALADase) enzyme activity, thereby effecting neuronal activities, inhibiting angiogenesis and treating glutamate abnormalities, compulsive disorders, prostate  
25 diseases, pain and diabetic neuropathy.

More specifically, the present invention relates to a compound of formula I:



or a pharmaceutically acceptable equivalent, wherein:

35 either J and K are taken together with one or more additional atoms independently selected from the group

consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 3-7 membered saturated or unsaturated heterocyclic or carbocyclic ring, and L is -CH,

or J, K, and L are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 4-8 membered saturated or unsaturated, mono-, bi-, or tricyclic, hetero- or carbocyclic ring structure;

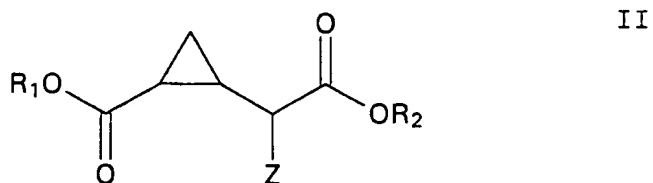
10 Z is a metal chelating group;

R<sub>1</sub> and R<sub>2</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

20 In a preferred embodiment of formula I, R<sub>1</sub> and R<sub>2</sub> are each hydrogen.

A preferred embodiment of the present invention relates to a compound of formula II:



25

or a pharmaceutically acceptable equivalent, wherein:

Z is a metal chelating group;

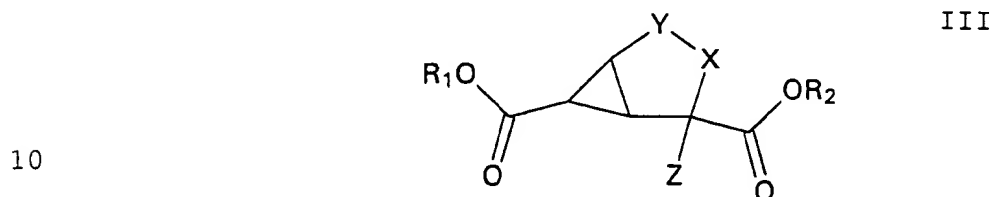
30 R<sub>1</sub> and R<sub>2</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

35 Ar is a carbocyclic or heterocyclic moiety which is

unsubstituted or substituted with one or more substituent(s).

In a preferred embodiment of formula II,  $R_1$  and  $R_2$  are each hydrogen.

5 Another preferred embodiment of the present invention relates to a compound of formula III:



or a pharmaceutically acceptable equivalent, wherein:

X and Y are independently selected from the group consisting of  $CH_2$ , O, NH, or S;

15 Z is a metal chelating group;

$R_1$  and  $R_2$  are independently hydrogen,  $C_1-C_6$  alkyl,  $C_2-C_6$  alkenyl,  $C_3-C_6$  cycloalkyl,  $C_5-C_7$  cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or  
20 more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

25 In a preferred embodiment of formula III,  $R_1$  and  $R_2$  are each hydrogen.

The present invention further relates to a method for treating a glutamate mediated disease, disorder, or condition in a mammal, comprising administering to said mammal an effective amount of a compound containing a  
30 metabotropic glutamate receptor ligand attached to a metal chelating group.

Finally, the present invention relates to a pharmaceutical composition comprising:

(i) an effective amount of a compound of formula I;  
35 and

(ii) a pharmaceutically acceptable carrier.

### INCORPORATION BY REFERENCE

5 In addition to other references generally known in the art, applicants have previously disclosed substantial data relating to the relationship between glutamate and various glutamate abnormalities, and the effectiveness of NAALADase inhibitors generally with regard to treating ischemic  
10 insult, diminished neurological function, alcohol dependence, nicotine dependence, cancer cell and tumor growth, angiogenesis, cocaine dependence, diabetes, pain, and hyperalgesia. Applicants hereby incorporate by reference, as though set forth herein in full, such figures and discussions from U.S. Patents Nos. 5,672,592,  
15 5,795,877, 5,863,536, 5,880,112 and 5,902,817, allowed U.S. Patent Applications Nos. 08/825,997, 08/833,628, 08/842,360 and 08/899,319 for which the issue fees have been paid, and International Publications Nos. WO 97/48399, WO 97/48400, WO 97/48409 and WO 98/53812. It would be expected that the  
20 compounds of the present invention would be effective as NAALADase inhibitors, and as such would have the same uses as the aforementioned compounds.

### 25 DETAILED DESCRIPTION OF THE INVENTION

#### Definitions

"Attached" refers to bonding, linkage, coupling, or other molecular association between atoms or molecules which results in a stable chemical entity.

30 "Chelate" or "chelation compound" refers to a coordination compound in which a central metal ion, such as  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Zn}^{2+}$ ,  $\text{Mg}^{2+}$ ,  $\text{Fe}^{2+}$ ,  $\text{Fe}^{3+}$ , or  $\text{Al}^{3+}$ , is attached by coordinate links to one or more nonmetal atoms in the same molecule, called chelating agents or ligands.

Further, a bidentate or polydentate chelating agent forms a heterocyclic ring with the central metal atom as part of each ring.

"Chelating agent," "chelating ligand," or "chelator" refers to any group which can chelate a metal. A chelating ligand offering one group for attachment to the metal is termed monodentate; two groups, bidentate; three or more groups, polydentate. A chelating ligand may attach to the metal atom by covalent or ionic bond(s). Many compounds, too numerous to fully enumerate here, can act as chelating ligands; common chelating ligands include, but are not limited to, derivatives of amines (e.g. ethylenediamine), aldehydes and ketones, carboxylic acids (e.g. ethylenediaminetetraacetic acid (EDTA)), sulfonyl- and mercapto-derivative groups, phosphoryls and other phosphorus derivatives, hydroxamic acid derivatives, and various combinations thereof.

"Derivative" refers to a substance produced from another substance either directly or by modification or partial substitution.

"Effective amount" refers to the amount required to produce the desired effect. "Therapeutically effective amount" refers to the amount required to inhibit NAALADase enzyme activity, effect neuronal activity, inhibit angiogenesis, and/or treat glutamate abnormality, compulsive disorder, prostate disease, pain and/or diabetic neuropathy.

"Electromagnetic radiation" includes without limitation radiation having the wavelength of  $10^{-20}$  to  $10^0$  meters. Preferred embodiments of the present invention employ the electromagnetic radiation of gamma-radiation ( $10^{-20}$  to  $10^{-13}$  m) x-ray radiation ( $10^{-11}$  to  $10^{-9}$  m), ultraviolet light (10 nm to 400 nm), visible light (400 nm to 700 nm), infrared radiation (700 nm to 1.0 mm), and microwave radiation (1 mm to 30 cm).

"Isosteres" refer to elements, molecules, or ions having different molecular formulae but exhibiting similar or identical physical properties. For example, tetrazole is an isostere of carboxylic acid because it mimics the properties of carboxylic acid even though they both have very different molecular formulae. Typically, two isosteric molecules have similar or identical volumes and shapes. Ideally, isosteric compounds should be isomorphic and able to co-crystallize. Among the other physical properties that isosteric compounds usually share are boiling point, density, viscosity and thermal conductivity. However, certain properties are usually different: dipolar moments, polarity, polarization, size and shape since the external orbitals may be hybridized differently. The term "isosteres" encompass "bioisosteres".

"Bioisosteres" are isosteres which, in addition to their physical similarities, share some common biological properties. Typically, bioisosteres interact with the same recognition site or produce broadly similar biological effects.

"Carboxylic acid isosteres" include without limitation direct derivatives such as hydroxamic acids, acyl-cyanamides and acylsulfonamides; planar acidic heterocycles such as tetrazoles, mercaptoazoles, sulfinylazoles, sulfonylazoles, isoxazoles, isothiazoles, hydroxythiadiazoles and hydroxychromes; and nonplanar sulfur- or phosphorus-derived acidic functions such as phosphinates, phosphonates, phosphonamides, sulphonates, sulphonamides, and acylsulphonamides.

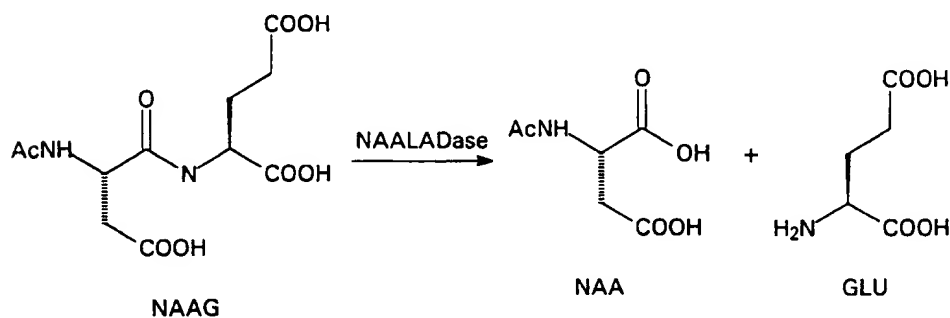
"Metabolite" refers to a substance produced by metabolism or by a metabolic process.

"NAAG" refers to N-acetyl-aspartyl-glutamate, an important peptide component of the brain, with levels comparable to the major inhibitor neurotransmitter gamma-aminobutyric acid (GABA). NAAG is neuron-specific, present

in synaptic vesicles and released upon neuronal stimulation in several systems presumed to be glutamatergic. Studies suggest that NAAG may function as a neurotransmitter and/or neuromodulator in the central nervous system, or as a precursor of the neurotransmitter glutamate. In addition, NAAG is an agonist at group II metabotropic glutamate receptors, specifically mGluR3 receptors; when attached to a moiety capable of inhibiting NAALADase, it is expected that metabotropic glutamate receptor ligands will provide potent and specific NAALADase inhibitors.

"NAALADase" refers to N-acetylated  $\alpha$ -linked acidic dipeptidase, a membrane bound metallopeptidase which catabolizes NAAG to N-acetylaspartate ("NAA") and glutamate ("GLU"):

**Catabolism of NAAG by NAALADase**



NAALADase has recently been assigned to the M28 peptidase family and is also called prostate specific membrane antigen (PSM) or human glutamate carboxypeptidase II (GCP II), EC number 3.4.17.21. It is believed that NAALADase is a co-catalytic zinc/zinc metallopeptidase. NAALADase shows a high affinity for NAAG with a  $K_m$  of 540 nM. If NAAG is a bioactive peptide, then NAALADase may serve to inactivate NAAG'S synaptic action. Alternatively, if NAAG functions as a precursor for glutamate, the primary function of NAALADase may be to regulate synaptic glutamate availability.

"Pharmaceutically acceptable carrier" refers to any carrier, diluent, excipient, suspending agent, lubricating

agent, adjuvant, vehicle, delivery system, emulsifier, disintegrant, absorbent, preservative, surfactant, colorant, flavorant, or sweetener. For these purposes, the compounds of the present invention may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally, vaginally or via an implanted reservoir in dosage formulations containing conventional non-toxic pharmaceutically-acceptable carriers, adjuvants and vehicles. The term parenteral as used herein includes subcutaneous, intravenous, intramuscular, intraperitoneal, intrathecal, intraventricular, intrasternal, and intracranial injection or infusion techniques.

"Pharmaceutically acceptable equivalent" includes, without limitation, pharmaceutically acceptable salts, hydrates, metabolites, prodrugs, and isosteres thereof. Many pharmaceutically acceptable equivalents are expected to have the same or similar *in vitro* or *in vivo* activity as the compounds of the invention.

"Pharmaceutically acceptable salt" refers to a salt of the inventive compounds which possesses the desired pharmacological activity and which is neither biologically nor otherwise undesirable. The salt can be formed with inorganic acids such as acetate, adipate, alginate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, glucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, thiocyanate, tosylate and undecanoate. Examples of a base salt include ammonium salts, alkali metal salts such as sodium and potassium salts, alkaline earth metal salts such as calcium and magnesium salts, salts with organic bases such as



dicyclohexylamine salts, N-methyl-D-glucamine, and salts with amino acids such as arginine and lysine. The basic nitrogen-containing groups can be quarternized with agents including lower alkyl halides such as methyl, ethyl, propyl and butyl chlorides, bromides and iodides; dialkyl sulfates such as dimethyl, diethyl, dibutyl and diamyl sulfates; long chain halides such as decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides; and aralkyl halides such as benzyl and phenethyl bromides.

"Pharmaceutically acceptable prodrug" refers to a derivative of the inventive compounds which undergoes biotransformation prior to exhibiting its pharmacological effect(s). A prodrug form is one which is not in an active form of the molecule as administered, but which becomes therapeutically active after some *in vivo* activity or biotransformation, such as metabolism, for example, enzymatic or hydrolytic cleavage. The prodrug is formulated with the objective(s) of improved chemical stability, improved patient acceptance and compliance, improved bioavailability, prolonged duration of action, improved organ selectivity, improved formulation (e.g., increased hydrosolubility), and/or decreased side effects (e.g., toxicity). The prodrug can be readily prepared from the inventive compounds using methods known in the art, such as those described by *Burger's Medicinal Chemistry and Drug Chemistry*, Fifth Ed., Vol. 1, pp. 172-178, 949-982 (1995). For example, the inventive compounds can be transformed into prodrugs by converting one or more of the hydroxy or carboxy groups into esters. Preferred prodrugs of the present invention include compounds of formula I, II, or III, where  $R_1$ ,  $R_2$ , or both are independently non-hydrogen moieties.

"Radiosensitizer" refers to a low molecular weight compound administered to animals in therapeutically effective amounts to promote the treatment of diseases

which are treatable with electromagnetic radiation. Diseases which are treatable with electromagnetic radiation include neoplastic diseases, benign and malignant tumors, and cancerous cells. Electromagnetic radiation treatment  
5 of other diseases not listed herein are also contemplated by the present invention.

"Alkyl" refers to a branched or unbranched saturated hydrocarbon chain comprising a designated number of carbon atoms. For example,  $C_1$ - $C_6$  straight or branched alkyl  
10 hydrocarbon chain contains 1 to 6 carbon atoms, and includes but is not limited to substituents such as methyl, ethyl, propyl, iso-propyl, butyl, iso-butyl, tert-butyl, n-pentyl, n-hexyl, and the like, unless otherwise indicated.

"Alkenyl" refers to a branched or unbranched unsaturated hydrocarbon chain comprising a designated  
15 number of carbon atoms. For example,  $C_2$ - $C_6$  straight or branched alkenyl hydrocarbon chain contains 2 to 6 carbon atoms having at least one double bond, and includes but is not limited to substituents such as ethenyl, propenyl, iso-propenyl, butenyl, iso-butenyl, tert-butenyl, n-pentenyl,  
20 n-hexenyl, and the like, unless otherwise indicated.

"Alkoxy" refers to the group -OR wherein R is alkyl as herein defined. Preferably, R is a branched or unbranched saturated hydrocarbon chain containing 1 to 6 carbon atoms.

"Aryl" or "aromatic" refers to an aromatic carbocyclic or heterocyclic group having a single ring, for example a phenyl ring; multiple rings, for example biphenyl; or multiple condensed rings in which at least one ring is aromatic, for example naphthyl, 1,2,3,4-tetrahydronaphthyl,  
30 anthryl, or phenanthryl, which can be unsubstituted or substituted with one or more other substituents as defined above. The substituents attached to a phenyl ring portion of an aryl moiety in the compounds of the invention may be configured in the ortho-, meta-, or para- orientations,  
35 with the para- orientation being preferred.

"Carbocycle" or "Carbocyclic moiety" refers to an organic cyclic moiety in which the cyclic skeleton is comprised of only carbon atoms, whereas the term "heterocycle" or "heterocyclic" refers to an organic cyclic moiety in which the cyclic skeleton contains one or more heteroatoms selected from nitrogen, oxygen, or sulfur, and which may or may not include carbon atoms. The term "carbocycle" refers to a carbocyclic moiety containing the indicated number of carbon atoms. The term "C<sub>3</sub>-C<sub>8</sub> cycloalkyl", therefore, refers to an organic cyclic substituent in which three to eight carbon atoms form a three, four, five, six, seven, or eight-membered ring, including, for example, a cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, or cyclooctyl ring.

"Heterocycle" or "heterocyclic moiety" refers to a saturated, unsaturated or aromatic carbocyclic group having a single ring, multiple fused rings (for example, bicyclic, tricyclic, or other similar bridged ring systems or substituents), or multiple condensed rings, and having at least one hetero atom such as nitrogen, oxygen, or sulfur within at least one of the rings. This term also includes "Heteroaryl" which refers to a heterocycle in which at least one ring is aromatic. Any heterocyclic or heteroaryl group can be unsubstituted or optionally substituted with one or more groups, as defined above.

"Inhibition", in the context of enzymes, refers to reversible enzyme inhibition such as competitive, uncompetitive and non-competitive inhibition. Competitive, uncompetitive and non-competitive inhibition can be distinguished by the effects of an inhibitor on the reaction kinetics of an enzyme. Competitive inhibition occurs when the inhibitor combines reversibly with the enzyme in such a way that it competes with a normal substrate for binding at the active site. The affinity between the inhibitor and the enzyme may be measured by the

inhibitor constant,  $K_i$ , which is defined as:

$$K_i = \frac{[E][I]}{[EI]}$$

wherein [E] is the concentration of the enzyme, [I] is the concentration of the inhibitor, and [EI] is the concentration of the enzyme-inhibitor complex formed by the reaction of the enzyme with the inhibitor. Unless otherwise specified,  $K_i$  as used herein refers to the affinity between the inventive compounds and NAALADase. "IC<sub>50</sub>" is a related term used to define the concentration or amount of a compound which is required to cause a 50% inhibition of the target enzyme. "NAALADase inhibitor" refers to any compound which inhibits NAALADase enzyme activity.

"Enantiomers" are a pair of stereoisomers that are non-superimposable mirror images of each other. "Enantiomer-enriched" refers to a mixture in which one enantiomer predominates.

"Isomers" refer to compounds having the same number and kind of atoms, and hence the same molecular weight, but differing in respect to the arrangement or configuration of the atoms.

"Optical isomers" refer to either of two kinds of stereoisomers. One kind is represented by mirror-image structures called enantiomers, which result from the presence of one or more asymmetric carbon atoms in the compound (glyceraldehyde, lactic acid, sugars, tartaric acid, amino acids). The other kind is exemplified by diastereoisomers, which are not mirror images. These occur in compounds having two or more asymmetric carbon atoms; thus, such compounds have  $2^n$  optical isomers, where  $n$  is the number of asymmetric carbon atoms.

"Stereoisomers" are isomers that differ only in the arrangement of the atoms in space. "Diastereoisomers" are stereoisomers which are not mirror images of each other.

5 "Racemic mixture" means a mixture containing equal parts of individual enantiomers. "Non-racemic mixture" is a mixture containing unequal parts of individual enantiomers or stereoisomers.

"Angiogenesis" refers to the process whereby new capillaries are formed.

10 "Angiogenesis-dependent disease" includes without limitation cancer, rheumatoid arthritis, cardiovascular disease, neovascular diseases of the eye, peripheral vascular disorders, and dermatologic ulcers. "Inhibition" of angiogenesis may be measured by many parameters in  
15 accordance with the present invention and, for instance, may be assessed by delayed appearance of neovascular structures, slowed development of neovascular structures, decreased occurrence of neovascular structures, slowed or decreased severity of angiogenesis-dependent disease  
20 effects, arrested angiogenic growth, or regression of previous angiogenic growth. In the extreme, complete inhibition is referred to herein as prevention. In relation to angiogenesis or angiogenic growth, "prevention" refers to no substantial angiogenesis or angiogenic growth  
25 if none had previously occurred, or no substantial further angiogenesis or angiogenic growth if growth had previously occurred.

"Animal" refers to a living organism having sensation and the power of voluntary movement, and which requires for  
30 its existence oxygen and organic food. Examples include, without limitation, a mammal such as a member of the human, equine, porcine, bovine, murine, canine, or feline species. In the case of a human, an "animal" may also be referred to as a "patient".

35 "Anxiety" includes without limitation the unpleasant

emotion state consisting of psychophysiological responses to anticipation of unreal or imagined danger, ostensibly resulting from unrecognized intrapsychic conflict. Physiological concomitants include increased heart rate, altered respiration rate, sweating, trembling, weakness, and fatigue; psychological concomitants include feelings of impending danger, powerlessness, apprehension, and tension. Dorland's Illustrated Medical Dictionary, W.B. Saunders Co., 27th ed. (1988).

"Anxiety Disorder" includes without limitation mental disorders in which anxiety and avoidance behavior predominate. Dorland's Illustrated Medical Dictionary, W.B. Saunders Co., 27th ed. (1988). Examples include without limitation panic attack, agoraphobia, panic disorder, acute stress disorder, chronic stress disorder, specific phobia, simple phobia, social phobia, substance induced anxiety disorder, organic anxiety disorder, obsessive compulsive disorder, post-traumatic stress disorder, generalized anxiety disorder, and anxiety disorder NOS. Other anxiety disorders are characterized in Diagnostic and Statistical Manual of Mental Disorders (American Psychiatric Association 4th ed. 1994). The skilled artisan will recognize that there are alternative nomenclatures, nosologies, and classification systems for pathologic psychological conditions and that these systems evolve with medical scientific progress.

"Attention Deficit Disorder" refers to a disorder characterized by developmentally inappropriate inattention and impulsiveness, with or without hyperactivity. Inattention means a failure to finish tasks started, easily distracted, seeming lack of attention, and difficulty concentrating on tasks requiring sustained attention. Impulsiveness means acting before thinking, difficulty taking turns, problems organizing work, and constant shifting from one activity to another. Hyperactivity means

difficulty staying seated and sitting still, and running or climbing excessively.

"Cancer" includes without limitation ACTH-producing tumors, acute lymphocytic leukemia, acute nonlymphocytic leukemia, cancer of the adrenal cortex, bladder cancer, brain cancer, breast cancer, cervix cancer, chronic lymphocytic leukemia, chronic myelocytic leukemia, colorectal cancer, cutaneous T-cell lymphoma, endometrial cancer, esophageal cancer, Ewing's sarcoma, gallbladder cancer, hairy cell leukemia, head and neck cancer, Hodgkin's lymphoma, Kaposi's sarcoma, kidney cancer, liver cancer, lung cancer (small and/or non-small cell), malignant peritoneal effusion, malignant pleural effusion, melanoma, mesothelioma, multiple myeloma, neuroblastoma, non-Hodgkin's lymphoma, osteosarcoma, ovary cancer, ovary (germ cell) cancer, pancreatic cancer, penis cancer, prostate cancer, retinoblastoma, skin cancer, soft-tissue sarcoma, squamous cell carcinomas, stomach cancer, testicular cancer, thyroid cancer, trophoblastic neoplasms, cancer of the uterus, vaginal cancer, cancer of the vulva, and Wilm's tumor.

"Compulsive disorder" refers to any disorder characterized by irresistible impulsive behavior. Examples of compulsive disorders include without limitation drug dependence, eating disorders, pathological gambling, ADD and Tourette's syndrome.

"Demyelinating diseases" refers to diseases of nerve tissue involving damage to or removal of the myelin sheath naturally surrounding such tissue. Such demyelinating diseases include, for example and without limitation, multiple sclerosis and peripheral demyelinating disease such as peripheral neuropathies and Charcot-Marie-Tooth disease.

"Disease" refers to any deviation from or interruption of the normal structure or function of any part, organ, or

system (or combination thereof) of the body that is manifested by a characteristic set of symptoms and signs and whose etiology, pathology, and prognosis may be known or unknown. Dorland's Illustrated Medical Dictionary,  
5 (W.B. Saunders Co. 27th ed. 1988).

"Disorder" refers to any derangement or abnormality of function; a morbid physical or mental state. Dorland's Illustrated Medical Dictionary, (W.B. Saunders Co. 27th ed. 1988).

10 "Drug dependence" refers to a psychologic addiction or a physical tolerance to a drug. Tolerance means a need to increase the dose progressively in order to produce the effect originally achieved by smaller amounts.

"Eating disorder" refers to compulsive overeating,  
15 obesity or severe obesity. Obesity means body weight of 20% over standard height-weight tables. Severe obesity means over 100% overweight.

"Glutamate abnormality" refers to any "disease, disorder, or condition" in which glutamate is implicated, including pathological conditions involving elevated levels  
20 of glutamate. Examples of glutamate abnormalities include, without limitation, spinal cord injury, epilepsy, stroke, Alzheimer's disease, Parkinson's disease, Amyotrophic Lateral Sclerosis (ALS), Huntington's disease,  
25 schizophrenia, pain, ischemia, peripheral neuropathy (including but not limited to diabetic neuropathy), traumatic brain injury, neuronal insult, inflammatory diseases, anxiety, anxiety disorders, memory impairment, and compulsive disorders.

30 "Glutamate modulator" refers to any composition of matter which alone or in combination with another agent affects the level of glutamate in a mammal.

"Ischemia" refers to localized tissue anemia due to obstruction of the inflow of arterial blood. Global  
35 ischemia occurs when blood flow ceases for a period of



time, such as may result from cardiac arrest. Focal ischemia occurs when a portion of the body, such as the brain, is deprived of its normal blood supply, such as may result from thromboembolytic occlusion of a cerebral vessel, traumatic head injury, edema or brain tumor. Even if transient, both global and focal ischemia can produce widespread neuronal damage. Although nerve tissue damage occurs over hours or even days following the onset of ischemia, some permanent nerve tissue damage may develop in the initial minutes following cessation of blood flow to the brain. Much of this damage is attributed to glutamate toxicity and secondary consequences of reperfusion of the tissue, such as the release of vasoactive products by damaged endothelium, and the release of cytotoxic products, such as free radicals and leukotrienes, by the damaged tissue.

"Memory impairment" refers to a diminished mental registration, retention or recall of past experiences, knowledge, ideas, sensations, thoughts or impressions. Memory impairment may affect short and long-term information retention, facility with spatial relationships, memory (rehearsal) strategies, and verbal retrieval and production. Common causes of memory impairment are age, severe head trauma, brain anoxia or ischemia, alcoholic-nutritional diseases, drug intoxications and neurodegenerative diseases. For example, memory impairment is a common feature of neurodegenerative diseases such as Alzheimer's disease and senile dementia of the Alzheimer type. Memory impairment also occurs with other kinds of dementia such as multi-infarct dementia, a senile dementia caused by cerebrovascular deficiency, and the Lewy-body variant of Alzheimer's disease with or without association with Parkinson's disease. Creutzfeldt-Jakob disease is a rare dementia with which memory impairment is associated. It is a spongiform encephalopathy caused by the prion

protein; it may be transmitted from other sufferers or may arise from gene mutations. Loss of memory is also a common feature of brain-damaged patients. Brain damage may occur, for example, after a classical stroke or as a result of an anaesthetic accident, head trauma, hypoglycemia, carbon monoxide poisoning, lithium intoxication, vitamin (B<sub>1</sub>, thiamine and B<sub>12</sub>) deficiency, or excessive alcohol use. Korsakoff's amnesic psychosis is a rare disorder characterized by profound memory loss and confabulation, whereby the patient invents stories to conceal his or her memory loss. It is frequently associated with excessive alcohol intake. Memory impairment may furthermore be age-associated; the ability to recall information such as names, places and words seems to decrease with increasing age. Transient memory loss may also occur in patients, suffering from a major depressive disorder, after electroconvulsive therapy.

"Enhancing memory performance" refers to improving or increasing the mental faculty by which to register, retain or recall past experiences, knowledge, ideas, sensations, thoughts or impressions.

"Mental disorder" refers to any clinically significant behavioral or psychological syndrome characterized by the presence of distressing symptoms or significant impairment of functioning. Mental disorders are assumed to result from some psychological or organic dysfunction of the individual; the concept does not include disturbances that are essentially conflicts between the individual and society (social deviance).

"Metastasis" refers to "[t]he ability of cells of a cancer to disseminate and form new foci of growth at noncontiguous sites (i.e., to form metastases)." See Hill, R.P, Chapter 11, "Metastasis", pp. 178-195 in *The Basic Science of Oncology*, Tannock et al., Eds., McGraw-Hill, New York (1992), herein incorporated by reference. "The

transition from *in situ* tumor growth to metastatic disease is defined by the ability of tumor cells of the primary site to invade local tissues and to cross tissue barriers ... To initiate the metastatic process, carcinoma cells  
5 must first penetrate the epithelial basement membrane and then invade the interstitial stroma ... For distant metastases, intravasation requires tumor cell invasion of the subendothelial basement membrane that must also be negotiated during tumor cell extravasation ... The  
10 development of malignancy is also associated with tumor-induced angiogenesis [which] not only allows for expansion of the primary tumors, but also permits easy access to the vascular compartment due to defects in the basement membranes of newly formed vessels." See Aznavoorian et  
15 al., *Cancer* 71: 1368-1383 (1993), herein incorporated by reference.

"Nervous function" refers to the various functions of the nervous system, which among other things provide an awareness of the internal and external environments of the  
20 body, make possible voluntary and reflex activities between the various structural elements of the organism, and balance the organism's response to environmental changes.

"Nervous insult" refers to any damage to nervous tissue and any disability or death resulting therefrom.  
25 The cause of nervous insult may be metabolic, toxic, neurotoxic, iatrogenic, thermal or chemical, and includes without limitation ischemia, hypoxia, cerebrovascular accident, trauma, surgery, pressure, mass effect, hemorrhage, radiation, vasospasm, neurodegenerative  
30 disease, neurodegenerative process, infection, Parkinson's disease, ALS, myelination/demyelination processes, epilepsy, cognitive disorder, glutamate abnormality and secondary effects thereof. Currently, there is no known effective treatment for nervous tissue damage.

"Nervous tissue" refers to the various components that make up the nervous system, including without limitation neurons, neural support cells, glia, Schwann cells, vasculature contained within and supplying these structures, the central nervous system, the brain, the brain stem, the spinal cord, the junction of the central nervous system with the peripheral nervous system, the peripheral nervous system, and allied structures.

"Neuroprotective" refers to the effect of reducing, arresting or ameliorating nervous insult, and protecting, resuscitating or reviving nervous tissue which has suffered nervous insult.

"Pain" refers to localized sensations of discomfort, distress or agony, resulting from the stimulation of specialized nerve endings. It serves as a protective mechanism insofar as it induces the sufferer to remove or withdraw from the source. Dorland's Illustrated Medical Dictionary, (W.B. Saunders Co. 27th ed. 1988). Examples of pain include, without limitation, acute, chronic, cancer, burn, incisional, inflammatory, diabetic neuropathic and back pain.

"Pathological gambling" refers to a condition characterized by a preoccupation with gambling. Similar to psychoactive substance abuse, its effects include development of tolerance with a need to gamble progressively larger amounts of money, withdrawal symptoms, and continued gambling despite severe negative effects on family and occupation.

"Prostate disease" refers to any disease affecting the prostate. Examples of prostate disease include without limitation prostate cancer such as adenocarcinoma and metastatic cancers of the prostate; and conditions characterized by abnormal growth of prostatic epithelial cells such as benign prostatic hyperplasia.

"Schizophrenia" refers to a mental disorder or group of mental disorders characterized by disturbances in form and content of thought (loosening of associations, delusions, hallucinations), mood (blunted, flattened, inappropriate affect), sense of self and relationship to the external world (loss of ego boundaries, dereistic thinking, and autistic withdrawal), and behavior (bizarre, apparently purposeless, and stereotyped activity or inactivity). Examples of schizophrenia include, without limitation, acute, ambulatory, borderline, catatonic, childhood, disorganized, hebephrenic, latent, nuclear, paranoid, paraphrenic, prepsychotic, process, pseudoneurotic, pseudopsychopathic, reactive, residual, schizo-affective and undifferentiated schizophrenia. Dorland's Illustrated Medical Dictionary, (W.B. Saunders Co. 27th ed. 1988).

"Therapeutic window of opportunity" or "window" refers, in relation to stroke, to the maximal delay between the onset of ischemia and the initiation of efficacious therapy.

"Tourette's syndrome" refers to an autosomal multiple tic disorder characterized by compulsive swearing, multiple muscle tics and loud noises. Tics are brief, rapid, involuntary movements that can be simple or complex; they are stereotyped and repetitive, but not rhythmic. Simple tics, such as eye blinking, often begin as nervous mannerisms. Complex tics often resemble fragments of normal behavior.

"Treating" or "treatment" as used herein covers any treatment of a disease and/or condition in a mammal, particularly a human, and includes:

- (i) preventing a disease, disorder or condition from occurring in a mammal which may be predisposed to the disease, disorder and/or condition but has not yet been diagnosed as having it;
- (ii) inhibiting the disease, disorder or condition,

i.e., arresting its development; and

(iii) relieving the disease, disorder or condition, i.e., causing regression of the disease, disorder and/or condition.

5 In relation to drug dependence, "treating" includes administering a compound or composition of the present invention to suppress the psychologic addiction or physical tolerance to the drug of abuse, and/or relieve and/or prevent a withdrawal syndrome resulting from the drug  
10 dependence.

"Withdrawal syndrome" refers to a disorder characterized by untoward physical changes that occur when the drug is discontinued or when its effect is counteracted by a specific antagonist.

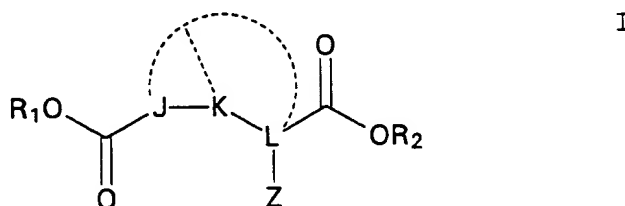
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# COMPOUNDS OF THE PRESENT INVENTION

## FORMULA I

The present invention relates to a compound of formula  
I:

20



25 or a pharmaceutically acceptable equivalent, wherein:

either J and K are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 3-7 membered saturated or  
30 unsaturated heterocyclic or carbocyclic ring, and L is -CH,

or J, K, and L are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 4-8 membered saturated or  
35 unsaturated, mono-, bi-, or tricyclic, hetero- or

carbocyclic ring structure;

Z is a metal chelating group;

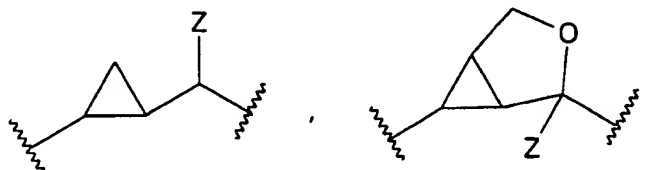
R<sub>1</sub> and R<sub>2</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein  
5 each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more  
10 substituent(s).

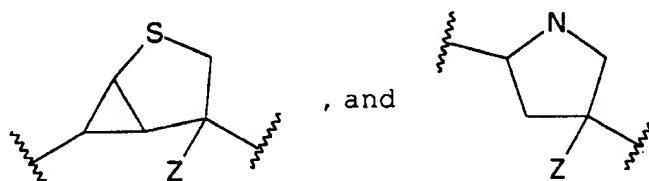
In a preferred embodiment of formula I, R<sub>1</sub> and R<sub>2</sub> are hydrogen.

Examples of preferred combinations of J, K, and L include, but are not limited to, the following:

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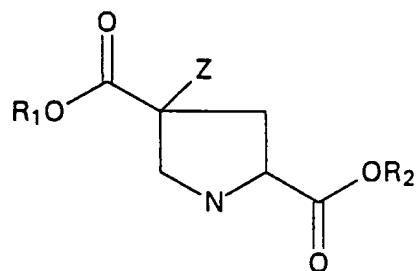
Possible substituents of said alkenyl, cycloalkyl, cycloalkenyl, and Ar include, without limitation, C<sub>1</sub>-C<sub>9</sub> straight or branched chain alkyl, C<sub>2</sub>-C<sub>9</sub> straight or branched chain alkenyl, C<sub>1</sub>-C<sub>9</sub> alkoxy, C<sub>2</sub>-C<sub>9</sub> alkenyloxy, phenoxy, benzyloxy, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, hydroxy, carboxy, carbonyl, amino, amido, cyano, isocyano, nitro, nitroso, nitrilo, isonitrilo, imino, azo, diazo, sulfonyl, sulfoxy, thio, thiocarbonyl, thiocyano, formamido, thioformamido, sulfhydryl, halo, haloalkyl, trifluoromethyl, and carbocyclic and heterocyclic moieties. Carbocyclic moieties include alicyclic and aromatic  
35 structures.

Examples of useful carbocyclic and heterocyclic moieties include, without limitation, phenyl, benzyl, naphthyl, indenyl, azulenyl, fluorenyl, anthracenyl, indolyl, isoindolyl, indolinyl, benzofuranyl, 5 benzothiophenyl, indazolyl, benzimidazolyl, benzthiazolyl, tetrahydrofuranyl, tetrahydropyranyl, pyridyl, pyrrolyl, pyrrolidinyl, pyridinyl, pyrimidinyl, purinyl, quinolinyl, isoquinolinyl, tetrahydroquinolinyl, quinoliziny, furyl, thiophenyl, imidazolyl, oxazolyl, benzoxazolyl, thiazolyl, 10 isoxazolyl, isotriazolyl, oxadiazolyl, triazolyl, thiadiazolyl, pyridazinyl, pyrimidinyl, pyrazinyl, triazinyl, trithianyl, indoliziny, pyrazolyl, pyrazolinyl, pyrazolidinyl, thienyl, tetrahydroisoquinolinyl, cinnolinyl, phthalazinyl, quinazolinyl, quinoxalinyl, 15 naphthyridinyl, pteridinyl, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, and phenoxazinyl.

Examples of useful metal chelating groups include, without limitation, mercapto derivatives, hydroxamic acid derivatives, phosphorus derivatives (particularly those of 20 the general formula  $X-P(O)(OH)-R$ , wherein R is as defined above for  $R_1$ ), carboxyl derivatives, N-carboxyalkyl derivatives, aldehydes, ketones, and combinations thereof. In particular, useful metal chelating groups include, without limitation, derivatives of dicarboxylic acids,  $\beta$ - 25 diketones,  $\alpha$ -hydroxycarboxylic acids, alkyl and aryl diamines,  $\alpha$ - and  $\beta$ -aminocarboxylates (including amino acid derivatives), thioethers, xanthates, dithiocarbamates, dithiocarboxylates, thioglycolates, thiols, and diphosphines. Preferred metal chelating ligands to be 30 substituted for Z in Formula I herein include, without limitation:





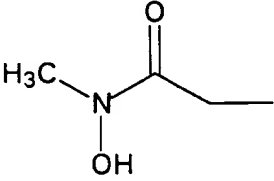
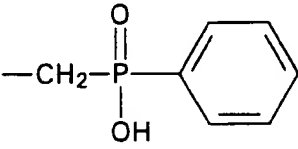
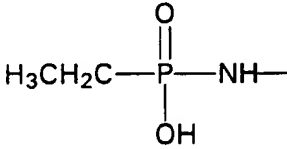
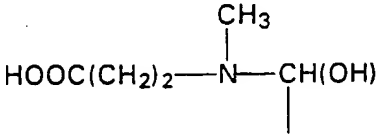
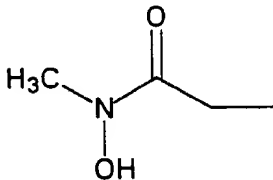
TABLE I

Compound	R <sub>1</sub>	R <sub>2</sub>	Z
1	H	H	-CH <sub>2</sub> P(O)(OH) <sub>2</sub>
2	H	H	
3	H	H	
4	H	H	
5	H	H	-CH <sub>2</sub> SH
6	H	H	

5

10

15

7	H	H	
8	H	H	$\text{HOOC}(\text{CH}_2)\text{—NH—CH}_2$ 
9	phenyl	H	$\text{—CH}_2\text{P}(\text{O})(\text{OH})_2$
10	$\text{—CH}_3$	$\text{—CH}_3$	$\text{—CH}_2\text{P}(\text{O})(\text{OH})\text{CH}_2\text{CH}_3$
11	H	$\text{—CH}_3$	
12	$\text{—CH}_2\text{CH}_3$	phenyl	$\text{—CH}_2\text{CH}_2\text{SH}$
13	cyclohexyl	$\text{—CH}_2\text{SH}$	
14	tri-fluoro-methyl	$\text{—NH}_2$	
15	pyridyl	benzyl	

Preferred compounds of formula I are selected from the group consisting of:

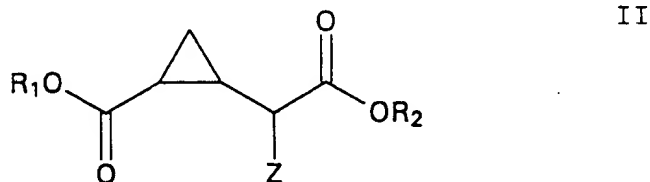
4-(phosphonomethyl)-2,4-pyrrolidine dicarboxylic acid (1);

4-[[hydroxy(phenyl)phosphinyl]methyl]-2,4-

pyrrolidinedicarboxylic acid (2);  
 4-[[hydroxy(phenylmethyl)phosphinyl]methyl]-2,4-  
 pyrrolidinedicarboxylic acid (3);  
 4-[[hydroxy(phenylethyl)phosphinyl]methyl]-2,4-  
 5 pyrrolidinedicarboxylic acid (4);  
 4-(sulfanylmethyl)-2,4-pyrrolidine dicarboxylic  
 acid (5); and  
 pharmaceutically acceptable equivalents thereof.

# **FORMULA II**

A preferred embodiment of the present invention relates to a compound of formula II:



or a pharmaceutically acceptable equivalent, wherein:

Z is a metal chelating group;

20  $R_1$  and  $R_2$  are independently hydrogen,  $C_1-C_9$  alkyl,  $C_2-C_9$  alkenyl,  $C_3-C_8$  cycloalkyl,  $C_5-C_7$  cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

25 Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

In a preferred embodiment of formula II,  $R_1$  and  $R_2$  are hydrogen.

30 Possible substituents of said alkenyl, cycloalkyl, cycloalkenyl, and Ar include, without limitation,  $C_1-C_9$  straight or branched chain alkyl,  $C_2-C_9$  straight or branched chain alkenyl,  $C_1-C_9$  alkoxy,  $C_2-C_9$  alkenyloxy, phenoxy, benzyloxy,  $C_3-C_8$  cycloalkyl,  $C_5-C_7$  cycloalkenyl, hydroxy, 35 carboxy, carbonyl, amino, amido, cyano, isocyano, nitro,

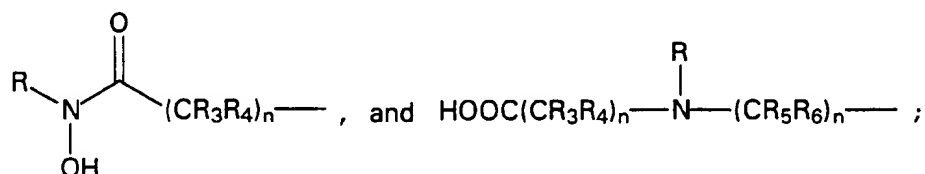
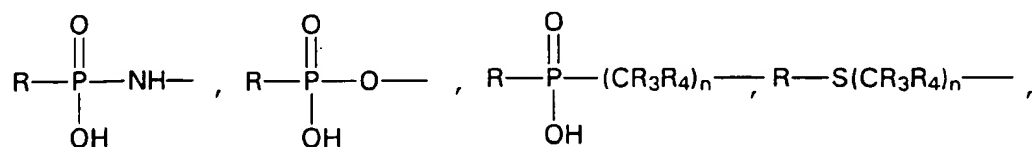
nitroso, nitrilo, isonitrilo, imino, azo, diazo, sulfonyl, sulfoxy, thio, thiocarbonyl, thiocyano, formanilido, thioformamido, sulfhydryl, halo, haloalkyl, trifluoromethyl, and carbocyclic and heterocyclic moieties.

5 Carbocyclic moieties include alicyclic and aromatic structures.

Examples of useful carbocyclic and heterocyclic moieties include, without limitation, phenyl, benzyl, naphthyl, indenyl, azulenyl, fluorenyl, anthracenyl, 10 indolyl, isoindolyl, indolinyl, benzofuranyl, benzothiophenyl, indazolyl, benzimidazolyl, benzthiazolyl, tetrahydrofuranyl, tetrahydropyranyl, pyridyl, pyrrolyl, pyrrolidinyl, pyridinyl, pyrimidinyl, purinyl, quinolinyl, isoquinolinyl, tetrahydroquinolinyl, quinoliziny, furyl, 15 thiophenyl, imidazolyl, oxazolyl, benzoxazolyl, thiazolyl, isoxazolyl, isotriazolyl, oxadiazolyl, triazolyl, thiadiazolyl, pyridazinyl, pyrimidinyl, pyrazinyl, triazinyl, trithianyl, indoliziny, pyrazolyl, pyrazolinyl, pyrazolidinyl, thienyl, tetrahydroisoquinolinyl, cinnolinyl, phthalazinyl, quinazolinyl, quinoxalinyl, 20 naphthyridinyl, pteridinyl, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, and phenoxazinyl.

Examples of useful metal chelating groups include, without limitation, mercapto derivatives, hydroxamic acid 25 derivatives, phosphorus derivatives (particularly those of the general formula  $X-P(O)(OH)-R$ , wherein R is as defined above for  $R_1$ ), carboxyl derivatives, N-carboxyalkyl derivatives, aldehydes, ketones, and combinations thereof. In particular, useful metal chelating groups include, 30 without limitation, derivatives of dicarboxylic acids,  $\beta$ -diketones,  $\alpha$ -hydroxycarboxylic acids, alkyl and aryl diamines,  $\alpha$ - and  $\beta$ -aminocarboxylates (including amino acid derivatives), thioethers, xanthates, dithiocarbamates, dithiocarboxylates, thioglycolates, thiols, and 35 diphosphines. Preferred metal chelating ligands to be

substituted for Z in Formula II include, without limitation:



wherein:

n is 0-3; and

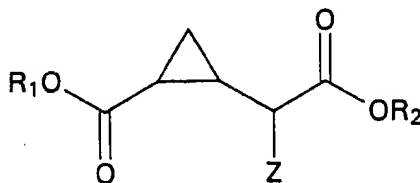
5 R, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>9</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

10 Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

Examples of compounds of formula II are shown in Table II:

15

**TABLE II**



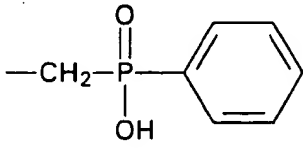
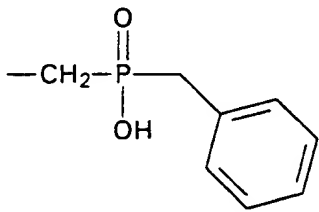
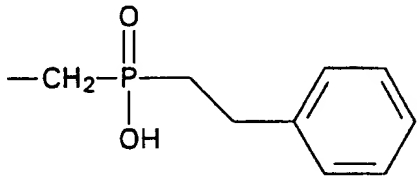
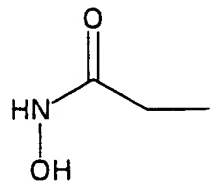
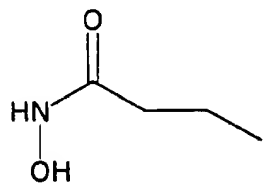
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Compound	R <sub>1</sub>	R <sub>2</sub>	Z
16	H	H	-CH <sub>2</sub> P(O)(OH) <sub>2</sub>
17	H	H	-CH <sub>2</sub> P(O)(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>
25 18	H	H	-CH <sub>2</sub> P(O)(OH)CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub>

5

10

15

19	H	H	
20	H	H	
21	H	H	
22	H	H	-CH <sub>2</sub> SH
23	H	H	-CH <sub>2</sub> CH <sub>2</sub> SH
24	H	H	-CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> SH
25	H	H	
26	H	H	
27	H	H	-NHCH <sub>2</sub> COOH
28	H	H	-CH <sub>2</sub> NHCH <sub>2</sub> COOH

Preferred compounds of formula II are selected from the group consisting of:

2-carboxy- $\alpha$ -(phosphonomethyl)-cyclopropaneacetic acid (16);

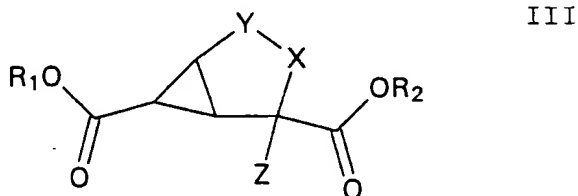
- 2-carboxy- $\alpha$ -[[hydroxypropylphosphinyl]methyl]-  
cyclopropaneacetic acid (17);  
2-carboxy- $\alpha$ -[[butylhydroxyphosphinyl]methyl]-  
cyclopropaneacetic acid (18);  
5 2-carboxy- $\alpha$ -[[hydroxyphenylphosphinyl]methyl]-  
cyclopropaneacetic acid (19);  
2-carboxy- $\alpha$ -[[hydroxy(phenylmethyl)phosphinyl]  
methyl]-cyclopropaneacetic acid (20);  
2-carboxy- $\alpha$ -[[hydroxy(2-phenylethyl)phosphinyl]  
10 methyl]-cyclopropaneacetic acid (21);  
2-carboxy- $\alpha$ -(mercaptoethyl)-cyclopropaneacetic  
acid (23);  
2-carboxy- $\alpha$ -(mercaptopropyl)-cyclopropaneacetic  
acid (24);  
15 2-carboxy- $\alpha$ -[2-(hydroxyamino)-2-oxoethyl]-  
cyclopropaneacetic acid (25);  
2-carboxy- $\alpha$ -[3-(hydroxyamino)-3-oxopropyl]-  
cyclopropaneacetic acid (26);  
2-carboxy- $\alpha$ -[(carboxymethyl)amino]-  
20 cyclopropaneacetic acid (27);  
2-carboxy- $\alpha$ -[[[(carboxymethyl)amino]methyl]-  
cyclopropaneacetic acid (28); and  
pharmaceutically acceptable equivalents.

25

**FORMULA III**

Another preferred embodiment of the present invention  
relates to a compound of formula III:

30



or a pharmaceutically acceptable equivalent, wherein:

- X and Y are independently selected from the group  
35 consisting of CH<sub>2</sub>, O, NH, or S;



Z is a metal chelating group;

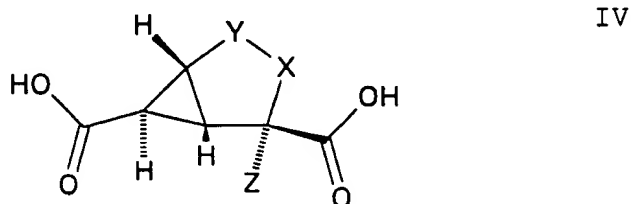
R<sub>1</sub> and R<sub>2</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein  
 5 is independently unsubstituted or substituted with one or more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

10 In a preferred embodiment of formula III, R<sub>1</sub> and R<sub>2</sub> are hydrogen.

In another preferred embodiment of formula III, the relative stereochemistry of the compound is of formula IV:

15



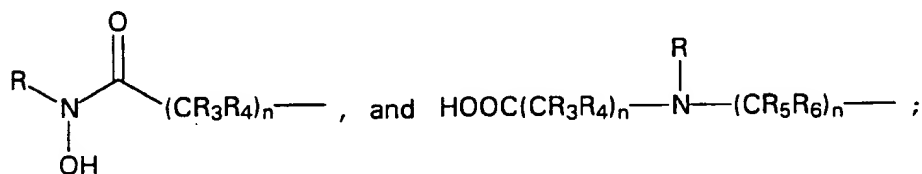
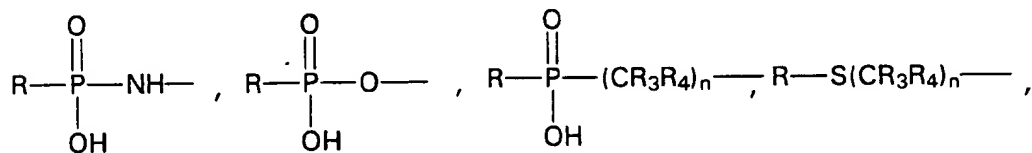
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Possible substituents of said alkenyl, cycloalkyl, cycloalkenyl, and Ar include, without limitation, C<sub>1</sub>-C<sub>9</sub> straight or branched chain alkyl, C<sub>2</sub>-C<sub>9</sub> straight or branched chain alkenyl, C<sub>1</sub>-C<sub>9</sub> alkoxy, C<sub>2</sub>-C<sub>9</sub> alkenyloxy, phenoxy, benzyloxy, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, hydroxy, carboxy, carbonyl, amino, amido, cyano, isocyano, nitro, nitroso, nitrilo, isonitrilo, imino, azo, diazo, sulfonyl, sulfoxy, thio, thiocarbonyl, thiocyno, formanilido, thioformamido, sulfhydryl, halo, haloalkyl, trifluoromethyl, and carbocyclic and heterocyclic moieties.  
 25  
 30 Carbocyclic moieties include alicyclic and aromatic structures.

Examples of useful carbocyclic and heterocyclic moieties include, without limitation, phenyl, benzyl, naphthyl, indenyl, azulenyl, fluorenyl, anthracenyl, indolyl, isoindolyl, indoliny, benzofuranyl,  
 35

benzothiophenyl, indazolyl, benzimidazolyl, benzthiazolyl, tetrahydrofuranyl, tetrahydropyranyl, pyridyl, pyrrolyl, pyrrolidinyl, pyridinyl, pyrimidinyl, purinyl, quinolinyl, isoquinolinyl, tetrahydroquinolinyl, quinolizinyl, furyl, thiophenyl, imidazolyl, oxazolyl, benzoxazolyl, thiazolyl, isoxazolyl, isotriazolyl, oxadiazolyl, triazolyl, thiadiazolyl, pyridazinyl, pyrimidinyl, pyrazinyl, triazinyl, trithianyl, indolizinyl, pyrazolyl, pyrazolinyl, pyrazolidinyl, thienyl, tetrahydroisoquinolinyl, cinnolinyl, phthalazinyl, quinazolinyl, quinoxalinyl, naphthyridinyl, pteridinyl, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, and phenoxazinyl.

Examples of useful metal chelating groups include, without limitation, mercapto derivatives, hydroxamic acid derivatives, phosphorus derivatives (particularly those of the general formula  $X-P(O)(OH)-R$ , wherein  $R$  is as defined above for  $R_1$ ), carboxyl derivatives, N-carboxyalkyl derivatives, aldehydes, ketones, and combinations thereof. In particular, useful metal chelating groups include, without limitation, derivatives of dicarboxylic acids,  $\beta$ -diketones,  $\alpha$ -hydroxycarboxylic acids, alkyl and aryl diamines,  $\alpha$ - and  $\beta$ -aminocarboxylates (including amino acid derivatives), thioethers, xanthates, dithiocarbamates, dithiocarboxylates, thioglycolates, thiols, and diphosphines. Preferred metal chelating ligands to be substituted for  $Z$  in Formula III herein include, without limitation:



wherein:

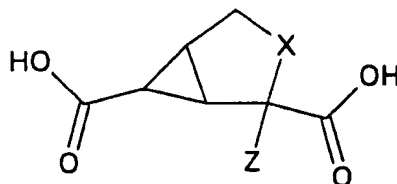
n is 0-3; and

R, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently hydrogen, C<sub>1</sub>-C<sub>3</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

Examples of compounds of formula III, where Y is carbon, and R<sub>1</sub> and R<sub>2</sub> are hydrogen, are shown in Table III:

**TABLE III**



Compound	X	Z
29	CH <sub>2</sub>	-CH <sub>2</sub> P(O)(OH) <sub>2</sub>
30	O	-CH <sub>2</sub> P(O)(OH)CH <sub>2</sub> CH <sub>3</sub>
31	CH <sub>2</sub>	

Preferred compounds of formula III are selected from the group consisting of:

2-(phosphonomethyl)-bicyclo[3.1.0]hexane-2,6-dicarboxylic acid (29);

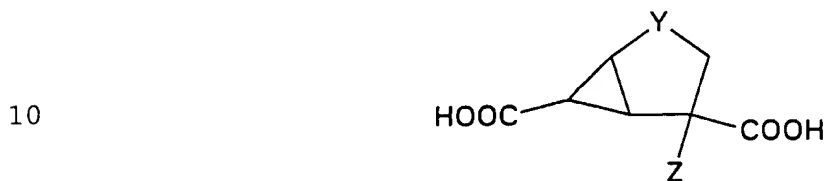
2-[[hydroxy(ethylphosphinyl)methyl]-3-oxabicyclo[3.1.0]hexane-2,6-dicarboxylic acid (30);

2-[[hydroxy(phenylmethyl)phosphinyl]methyl]-

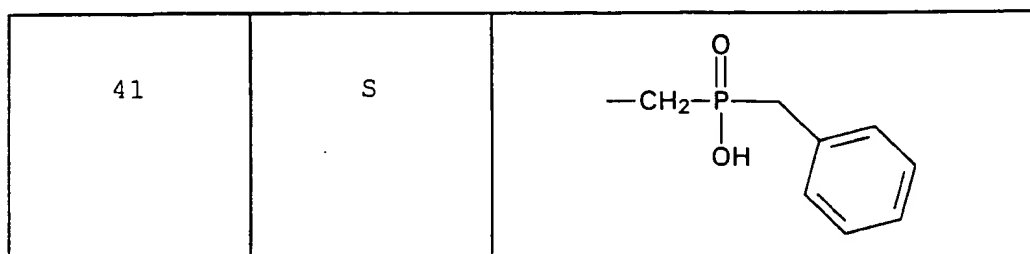
bicyclo[3.1.0]hexane-2,6-dicarboxylic acid (31);  
and  
pharmaceutically acceptable equivalents.

- 5 Examples of compounds of formula III, where X is carbon, and R<sub>1</sub> and R<sub>2</sub> are hydrogen, are shown in Table IV:

**TABLE IV**



Compound	Y	Z
15 32	O	$\text{H}_3\text{CH}_2\text{C}-\text{P}\begin{matrix} \text{O} \\ \parallel \\ \text{OH} \end{matrix}-\text{NH}-$
33	S	$\text{H}_3\text{CH}_2\text{C}-\text{P}\begin{matrix} \text{O} \\ \parallel \\ \text{OH} \end{matrix}-\text{NH}-$
34	O	$-\text{CH}_2\text{P}(\text{O})(\text{OH})_2$
35	S	$-\text{CH}_2\text{P}(\text{O})(\text{OH})_2$
20 36	O	$-\text{CH}_2\text{SH}$
37	S	$-\text{CH}_2\text{SH}$
38	O	$-\text{CH}_2\text{P}(\text{O})(\text{OH})\text{CH}_2\text{CH}_3$
39	S	$-\text{CH}_2\text{P}(\text{O})(\text{OH})\text{CH}_2\text{CH}_3$
25 40	O	$-\text{CH}_2-\text{P}\begin{matrix} \text{O} \\ \parallel \\ \text{OH} \end{matrix}-\text{CH}_2-\text{C}_6\text{H}_5$



5 Preferred compounds of formula III are selected from the group consisting of:

- 2-oxa-4-(ethylhydroxyphosphoryl)aminobicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 2-thia-4-(ethylhydroxyphosphoryl)aminobicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 10 2-oxa-4-(hydroxyphosphoryl)bicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 2-thia-4-(hydroxyphosphoryl)bicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 15 2-oxa-4-(methylsulfanyl)bicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 2-thia-4-(methylsulfanyl)bicyclo[3.1.0]hexane-4,6-dicarboxylate;
- 4-[[hydroxy(phenylmethyl)phosphinyl]methyl]-2-oxabicyclo[3.1.0]hexane-4,6-dicarboxylic acid;
- 20 and
- pharmaceutically acceptable equivalents thereof.

As already stated, the invention includes within its

25 scope pharmaceutically acceptable equivalents of the inventive compounds, such as pharmaceutically acceptable salts and prodrugs. Examples of preferred pharmaceutically acceptable salts are either those with inorganic bases, such as sodium, potassium, calcium and aluminum hydroxides,

30 or with organic bases, such as lysine, arginine, N-methyl-glucamine, triethylamine, triethanolamine, dibenzylamine, methylbenzylamine, di-(2-ethyl-hexyl)-amine,

piperidine, N-ethylpiperidine, N,N-diethylaminoethylamine, N-ethylmorpholine,  $\beta$ -phenethylamine, N-benzyl- $\beta$ -phenethylamine, N-benzyl-N,N-dimethylamine and other acceptable organic amines.

5 Preferred prodrugs include compounds of formulas I, II, and III, where  $R_1$  and  $R_2$  are non-hydrogen moieties.

Some compounds of the present invention possess one or more asymmetric carbon center(s) and are thus capable of existing in the form of optical isomers as well as in the form of racemic or non-racemic mixtures of optical isomers. The optical isomers can be obtained by resolution of the racemic mixtures according to conventional processes well known in the art, for example by formation of diastereoisomeric salts by treatment with an optically active acid or base. Examples of appropriate acids are tartaric, diacetyltartaric, dibenzoyltartaric, ditoluoyltartaric and camphorsulfonic acid and then separation of the mixture of diastereoisomers by crystallization followed by liberation of the optically active bases from these salts. A different process for separation of optical isomers involves the use of a chiral chromatography column optimally chosen to maximize the separation of the enantiomers. Still another available method involves synthesis of covalent diastereoisomeric molecules, for example, esters, amides, acetals, ketals, and the like, by reacting compounds used in the inventive methods and pharmaceutical compositions with an optically active acid in an activated form, an optically active diol or an optically active isocyanate. The synthesized diastereoisomers can be separated by conventional means such as chromatography, distillation, crystallization or sublimation, and then hydrolyzed to deliver the enantiomerically pure compound. In some cases hydrolysis to the parent optically active drug is not necessary prior to dosing the patient since the compound can behave as a

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prodrug. The optically active compounds of the present invention can likewise be obtained by utilizing optically active starting materials.

It is understood that the inventive compounds encompass optical isomers as well as racemic and non-racemic mixtures.

#### **PHARMACEUTICAL COMPOSITIONS OF THE PRESENT INVENTION**

The present invention relates to a pharmaceutical composition comprising:

(i) an effective amount of a compound of the present invention; and

(ii) a pharmaceutically acceptable carrier.

Preferred compounds of the present invention are set forth above.

Preferably, the compound of the present invention is present in an effective amount for inhibiting NAALADase enzyme activity, treating a glutamate abnormality, effecting a neuronal activity, treating a compulsive disorder, treating a prostate disease, or inhibiting angiogenesis in a mammal.

#### **METHODS OF THE PRESENT INVENTION**

##### **METHOD FOR INHIBITING NAALADASE ENZYME ACTIVITY**

The present invention relates to a method for inhibiting NAALADase enzyme activity in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

NAAG is the natural substrate for the NAALADase enzyme. NAAG is also an agonist at group II metabotropic glutamate receptors, specifically mGluR3 receptors. The majority of the metabotropic glutamate receptor ligands known are cyclized glutamate analogs. Thus, in a preferred embodiment, combining a metal chelating group capable of

interacting with the metal atom(s) at the active site of NAALADase with a metabotropic glutamate receptor ligand, preferably an mGluR3 receptor ligand, in the form of a cyclized glutamate unit, is expected to provide a potent and specific NAALADase inhibitor.

#### **METHOD FOR TREATING GLUTAMATE ABNORMALITY**

The present invention further relates to a method for treating a "glutamate abnormality" in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

Although not limited to any one particular theory, it is believed that the compounds of the present invention modulate levels of glutamate by acting on a storage form of glutamate which is hypothesized to be upstream from the effects mediated by the NMDA receptor.

#### **METHOD FOR TREATING COMPULSIVE DISORDER**

The present invention further relates to a method for treating a compulsive disorder, comprising administering to a patient in need of such treatment an effective amount of a compound of the present invention.

#### **METHOD FOR EFFECTING NEURONAL ACTIVITY**

The present invention further relates to a method for effecting a neuronal activity in a mammal, comprising administering to said mammal an effective amount of the compound of the present invention.

The neuronal activity that is effected by the inventive method may be selected from the group consisting of: stimulation of damaged neurons, promotion of neuronal regeneration, prevention of neurodegeneration and treatment of a neurological disorder.

Examples of neurological disorders that are treatable by the methods of the present invention include without



limitation: trigeminal neuralgia; glossopharyngeal neuralgia; Bell's Palsy; myasthenia gravis; muscular dystrophy; amyotrophic lateral sclerosis; progressive muscular atrophy; progressive bulbar inherited muscular atrophy; herniated, ruptured or prolapsed invertebrate disk syndromes; cervical spondylosis; plexus disorders; thoracic outlet destruction syndromes; peripheral neuropathies such as those caused by lead, dapsone, ticks, porphyria, or Guillain-Barré syndrome; Alzheimer's disease; and Parkinson's disease.

The inventive method is particularly useful for treating a neurological disorder selected from the group consisting of: peripheral neuropathy caused by physical injury or disease state, traumatic brain injury, physical damage to the spinal cord, stroke associated with brain damage, demyelinating diseases and neurological disorders relating to neurodegeneration. Examples of demyelinating diseases include multiple sclerosis and peripheral demyelinating disease such as peripheral neuropathies and Charcot-Marie Tooth disease. Examples of neurological disorders relating to neurodegeneration include Alzheimer's disease, Parkinson's disease, and amyotrophic lateral sclerosis (ALS).

#### **METHOD FOR TREATING PROSTATE DISEASE**

The present invention further relates to a method for treating a prostate disease in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

30

#### **METHOD FOR TREATING CANCER**

In addition to prostate cancer, other forms of cancer may be treated with the compounds of the present invention. The compounds of the present invention are particularly useful in treating cancer of tissues where NAALADase

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enzymes reside. Such tissues include the prostate as well as the brain, kidney and testis.

#### METHOD FOR TREATING STROKE

5       The present invention further relates to a method for treating stroke in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention more than 60 minutes following onset of stroke. Preferably, the compound is administered to said  
10       mammal more than 120 minutes following the onset of stroke.

#### METHOD FOR INHIBITING ANGIOGENESIS

It is expected that the compounds of the present invention can affect angiogenesis in tissues containing  
15       NAALADase. Previous research showed that NAALADase is enriched in synaptic plasma membranes and is primarily localized to neural and kidney tissue. NAALADase has also been found in the tissues of the prostate and testes. Additionally, previous findings have shown NAALADase to be  
20       present in neovasculature. Furthermore, as NAALADase continues to be discovered in other tissues of the body, NAALADase inhibitors most likely will also show efficacy in the inhibition of angiogenesis in those tissues.

Accordingly, the present invention further relates to  
25       a method for inhibiting angiogenesis in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

Angiogenesis may be necessary for fertility or metastasis of cancer tumors, or may be related to an  
30       angiogenic-dependent disease. Thus, the angiogenic-dependent diseases treatable by the inventive methods include without limitation rheumatoid arthritis, cardiovascular diseases, neovascular diseases of the eye, peripheral vascular disorders, and cancerous tumor growth,  
35       invasion, and metastasis.

#### METHOD FOR TREATING PAIN

The present invention further relates to a method for treating pain in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

The compounds of the present invention are particularly effective in blocking tolerance to morphine and reducing the amount of morphine necessary for treating pain. In a preferred embodiment, the compound of the present invention is administered in combination with morphine.

#### METHOD FOR TREATING DIABETIC NEUROPATHY

The present invention further relates to a method for treating diabetic neuropathy in a mammal, comprising administering to said mammal an effective amount of a compound of the present invention.

#### ROUTE OF ADMINISTRATION

The compounds of the present invention will generally be administered by means well known to the ordinarily skilled artisan to a patient in the form of a pharmaceutical formulation. Such formulations preferably include, in addition to the active agent, a physiologically acceptable carrier and/or diluent. In the methods of the present invention, the compounds may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally, vaginally or via an implanted reservoir in dosage formulations containing conventional non-toxic pharmaceutically-acceptable carriers, adjuvants and vehicles. The term parenteral as used herein includes subcutaneous, intravenous, intramuscular, intraperitoneal, intrathecal, intraventricular, intrasternal, intracranial or intraosseous injection and infusion techniques.

To be effective therapeutically as central nervous system targets, the compounds of the present invention should readily penetrate the blood-brain barrier when peripherally administered. Compounds which cannot  
5 penetrate the blood-brain barrier can be effectively administered by an intraventricular route.

#### DOSAGE

The compounds of the present invention may be  
10 administered by a single dose, multiple discrete doses or continuous infusion. Since the compounds are small, easily diffusible and relatively stable, they are well suited to continuous infusion. Pump means, particularly subcutaneous pump means, are preferred for continuous infusion.

15 Dose levels on the order of about 0.1 mg to about 10,000 mg of the active ingredient compound are useful in the treatment of the above conditions, with preferred levels being about 0.1 mg to about 1,000 mg. The specific dose level for any particular patient will vary depending  
20 upon a variety of factors, including the activity of the specific compound employed; the age, body weight, general health, sex and diet of the patient; the time of administration; the rate of excretion; drug combination; the severity of the particular disease being treated; and  
25 the form of administration. Typically, *in vitro* dosage-effect results provide useful guidance on the proper doses for patient administration. Studies in animal models are also helpful. The considerations for determining the proper dose levels are well known in the art.

30

#### ADMINISTRATION REGIMEN

For the methods of the present invention, any administration regimen well known to the ordinarily skilled artisan for regulating the timing and sequence of drug  
35 delivery can be used and repeated as necessary to effect

treatment. Such regimen may include pretreatment and/or co-administration with additional therapeutic agents.

#### COMBINATION WITH OTHER TREATMENTS

5 The compounds of the present invention may be used alone or in combination with other biologically active agent(s) for simultaneous, separate, or sequential use.

The biologically active agent may be selected from a wide variety of materials, including but not limited to  
10 steroids, for example hydrocortisomers such as methylprednisolone; anti-inflammatory or anti-immune drugs, such as methotrexate, azathioprine, cyclophosphamide or cyclosporin A; interferon- $\beta$ ; antibodies, such as anti-CD4 antibodies; agents which can reduce the risk of a  
15 second ischemic event, such as ticlopidine; chemotherapeutic agents; immunotherapeutic compositions; electromagnetic radiosensitizers; morphine for treating pain; or mixtures thereof.

The compounds of the present invention can be co-administered with one or more therapeutic agents either (i)  
20 together in a single formulation, or (ii) separately in individual formulations designed for optimal release rates of their respective active agent. Each formulation may contain from about 0.01% to about 99.99% by weight,  
25 preferably from about 3.5% to about 60% by weight, of a compound of the present invention, as well as one or more pharmaceutical excipients, such as wetting, emulsifying and pH buffering agents.

#### EXPERIMENTAL DATA

30 Applicants have previously disclosed substantial data relating to the relationship between glutamate and various glutamate abnormalities, and the effectiveness of NAALADase inhibitors. See "Incorporation by Reference" above for  
35 further details.

### EXAMPLES

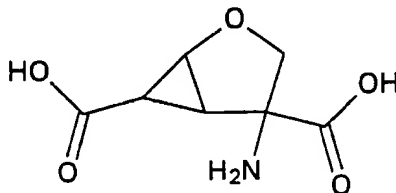
The following examples are illustrative of the present invention and are not intended to be limitations thereon. Unless otherwise indicated, all percentages are based upon 100% by weight of the final composition.

The compounds of the present invention possess one or more asymmetric center(s) and thus can be produced as mixtures (racemic and non-racemic) of stereoisomers, or as individual R- and S-stereoisomers. The individual stereoisomers may be obtained by using an optically active starting material, by resolving a racemic or non-racemic mixture of an intermediate at some appropriate stage of synthesis, or by resolving a compound of the present invention. It is understood that the compounds of the present invention encompass optical isomers, individual stereoisomers as well as mixtures (racemic and non-racemic) of stereoisomers.

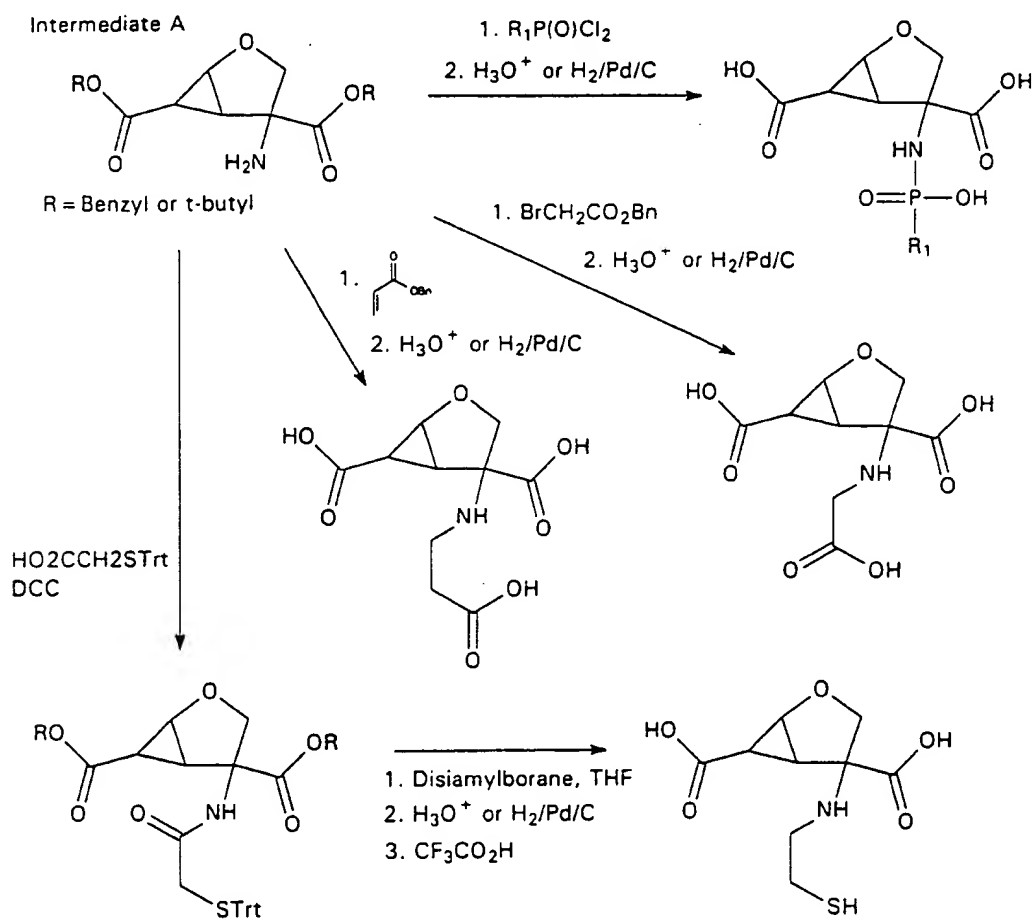
#### EXAMPLE 1

##### Synthesis of Compounds

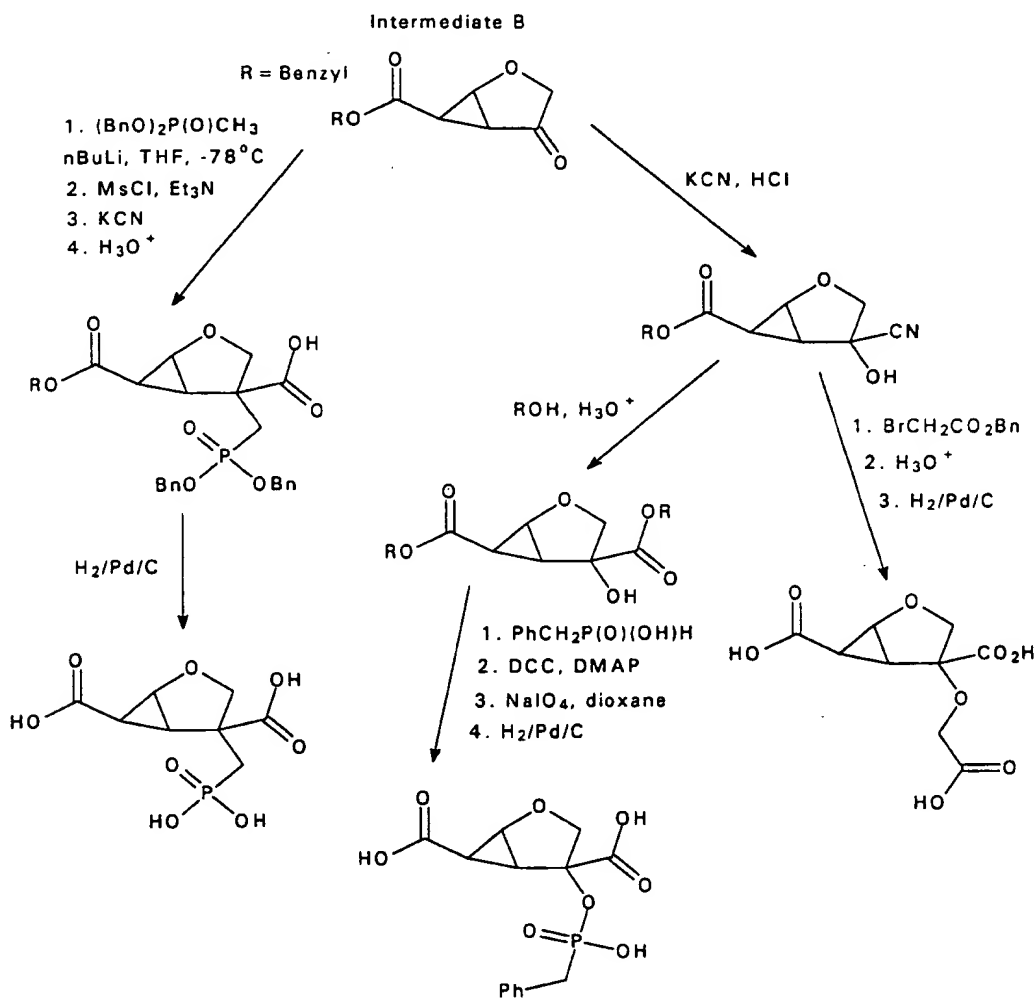
The compounds of the present invention can be readily prepared by a variety of standard techniques of organic chemistry known in the art. Precursor compounds can also be prepared by methods known in the art. For Example, the following intermediate has been described by Monn, et al., in *J. Med. Chem.* 42:1027-1040, 1999:



This intermediate can be converted to a protected acid using standard techniques of organic chemistry, to provide Intermediate A, which in turn can be converted to compounds of the invention by the pathways depicted in Scheme I:

SCHEME I

Further, Intermediate B can be prepared by the general method described by Monn, et al., in *J. Med. Chem.* 42:1027-1040, 1999, which in turn can be converted to compounds of the invention by the pathways depicted in Scheme II:

SCHEME IIEXAMPLE 2

5        A patient is suffering from a disease, disorder or condition described. The patient may then be administered an effective amount of a compound or a pharmaceutical composition of the present invention. It is expected that after such treatment, the patient would not suffer any

10      significant injury due to, would be protected from further injury due to, or would recover from the disease, disorder or condition.



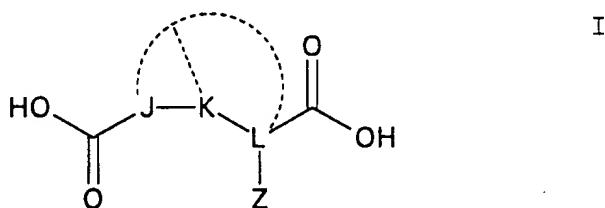
The invention being thus described, it will be obvious that the same may be varied in many ways. Such variations are not to be regarded as a departure from the spirit and scope of the invention and all such modifications are intended to be included within the scope of the following claims.

**WE CLAIM:**

1. A compound containing a metabotropic glutamate receptor ligand attached to a metal chelating group.

2. The compound of claim 1, wherein the metabotropic glutamate receptor is mGluR3.

3. The compound of claim 1, wherein the compound is of formula I:



or a pharmaceutically acceptable equivalent, wherein:

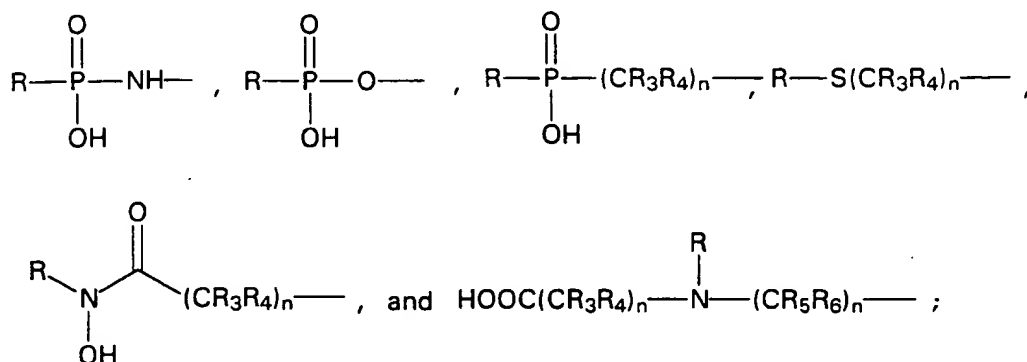
either J and K are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 3-7 membered saturated or unsaturated heterocyclic or carbocyclic ring, and L is -CH,

or J, K, and L are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 4-8 membered saturated or unsaturated, mono-, bi-, or tricyclic, hetero- or carbocyclic ring structure; and

Z is a metal chelating group.

4. The compound of claim 3, wherein said metal chelating group is selected from the group consisting of:

58



wherein:

n is 0-3; and

R, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

5. The compound of claim 4, wherein the compound is selected from the group consisting of:

4-(phosphonomethyl)-2,4-pyrrolidine dicarboxylic acid;

4-[[hydroxy(phenyl)phosphinyl]methyl]-2,4-pyrrolidinedicarboxylic acid;

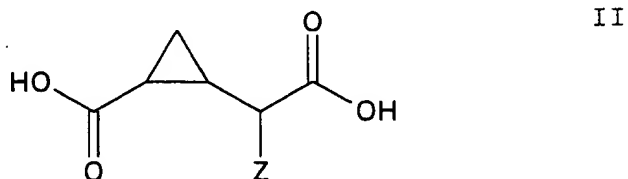
4-[[hydroxy(phenylmethyl)phosphinyl]methyl]-2,4-pyrrolidinecarboxylic acid;

4-[[hydroxy(phenylethyl)phosphinyl]methyl]-2,4-pyrrolidinedicarboxylic acid; and

4-(sulfanylmethyl)-2,4-pyrrolidine dicarboxylic acid; and

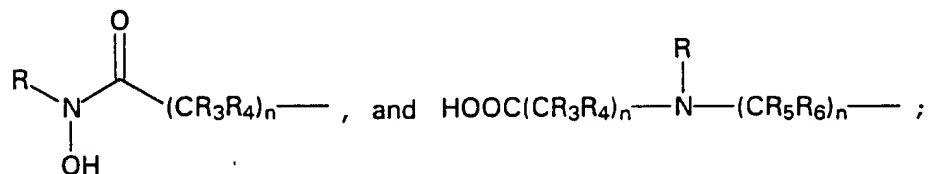
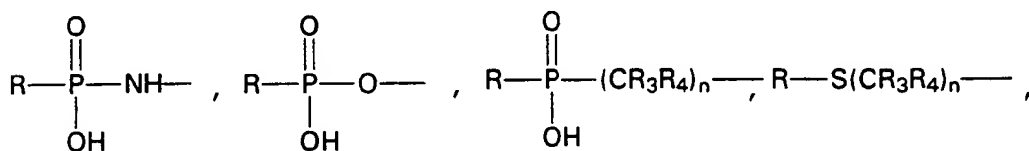
pharmaceutically acceptable equivalents.

6. The compound of claim 3, wherein the compound is of formula II:



or a pharmaceutically acceptable equivalent.

7. The compound of claim 6, wherein Z is selected from the group consisting of:



wherein:

n is 0-3; and

R, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

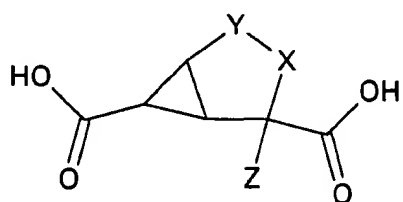
Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

8. The compound of claim 7, wherein said compound is selected from the group consisting of:

2-carboxy-α-(phosphonomethyl)-cyclopropaneacetic acid;

- 2-carboxy- $\alpha$ -[[hydroxypropylphosphinyl]methyl]-  
cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[[butylhydroxyphosphinyl]methyl]-  
cyclopropaneacetic acid;  
5 2-carboxy- $\alpha$ -[[hydroxyphenylphosphinyl]methyl]-  
cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[[hydroxy(phenylmethyl)phosphinyl]  
methyl]-cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[[hydroxy(2-phenylethyl)phosphinyl]  
10 methyl]-cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -(mercaptoethyl)-cyclopropaneacetic  
acid;  
2-carboxy- $\alpha$ -(mercaptopropyl)-cyclopropaneacetic  
acid;  
15 2-carboxy- $\alpha$ -[2-(hydroxyamino)-2-oxoethyl]-  
cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[3-(hydroxyamino)-3-oxopropyl]-  
cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[(carboxymethyl)amino]-  
20 cyclopropaneacetic acid;  
2-carboxy- $\alpha$ -[[ (carboxymethyl)amino]methyl]-  
cyclopropaneacetic acid; and  
pharmaceutically acceptable equivalents.

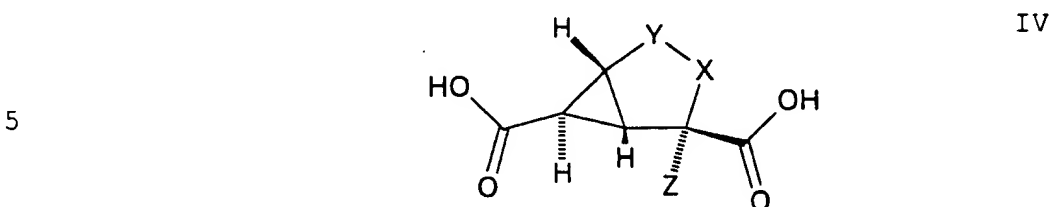
- 25 9. The compound of claim 3, wherein the compound is  
of formula III:



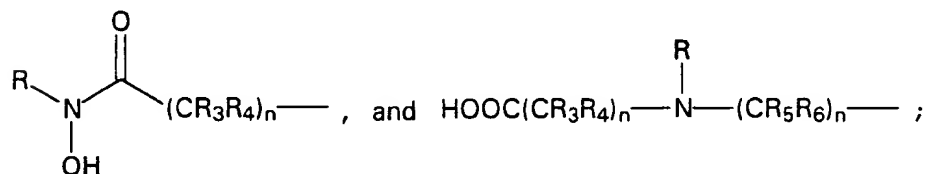
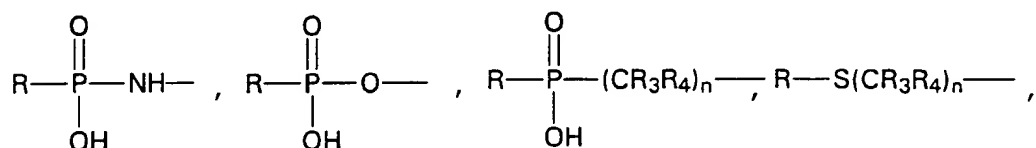
or a pharmaceutically acceptable equivalent, wherein:

X and Y are independently selected from the group  
consisting of CH<sub>2</sub>, O, NH, or S.

10. The compound of claim 9, wherein the relative stereochemistry of the compound is of formula IV:



10 11. The compound of claim 9, wherein Z is selected from the group consisting of:



wherein:

n is 0-3; and

15 R, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently hydrogen, C<sub>1</sub>-C<sub>9</sub> alkyl, C<sub>2</sub>-C<sub>9</sub> alkenyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, C<sub>5</sub>-C<sub>7</sub> cycloalkenyl, or Ar, wherein each said alkyl, alkenyl, cycloalkyl, cycloalkenyl, or Ar is independently unsubstituted or substituted with one or more substituent(s); and

20 Ar is a carbocyclic or heterocyclic moiety which is unsubstituted or substituted with one or more substituent(s).

12. The compound of claim 11, wherein said compound is selected from the group consisting of:

25 2-(phosphonomethyl)-bicyclo[3.1.0]hexane-2,6-dicarboxylic acid;

2-[[hydroxy(ethylphosphinyl)methyl]-3-oxabicyclo

[3.1.0]hexane-2,6-dicarboxylic acid;  
2-[[hydroxy(phenylmethyl)phosphinyl]methyl]-  
bicyclo[3.1.0]hexane-2,6-dicarboxylic acid;  
2-oxa-4-(ethylhydroxyphosphoryl)  
5 aminobicyclo[3.1.0]hexane-4,6-dicarboxylate;  
2-thia-4-(ethylhydroxyphosphoryl)  
aminobicyclo[3.1.0]hexane-4,6-dicarboxylate;  
2-oxa-4-(hydroxyphosphoryl)bicyclo[3.1.0]hexane-  
4,6-dicarboxylate;  
10 2-thia-4-(hydroxyphosphoryl)bicyclo[3.1.0]hexane-  
4,6-dicarboxylate;  
2-oxa-4-(methylsulfanyl)bicyclo[3.1.0]hexane-4,6-  
dicarboxylate;  
2-thia-4-(methylsulfanyl)bicyclo[3.1.0]hexane-  
15 4,6-dicarboxylate;  
4-[[hydroxy(phenylmethyl)phosphinyl]methyl]-2-  
oxabicyclo[3.1.0]hexane-4,6-dicarboxylic acid;  
and  
pharmaceutically acceptable equivalents.

20

13. The compound of claim 1, wherein said  
metabotropic glutamate receptor ligand is quisqualic acid.

25

14. A method for inhibiting NAALADase enzyme activity  
in a mammal, comprising administering to said mammal an  
effective amount of a compound containing a metabotropic  
glutamate receptor ligand attached to a metal chelating  
group.

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15. A method for treating a glutamate mediated  
disease, disorder, or condition in a mammal, comprising  
administering to said mammal an effective amount of a  
compound containing a metabotropic glutamate receptor  
ligand attached to a metal chelating group.

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16. The method of claim 15, wherein the glutamate mediated disease, disorder, or condition is selected from the group consisting of: stroke, Alzheimer's disease, Parkinson's disease, ALS, Huntington's disease, schizophrenia, ischemia, pain, diabetic neuropathy, trauma, nervous insult, anxiety, anxiety disorder, inflammatory diseases, and memory impairment.

17. The method of claim 16, wherein the glutamate mediated disease, disorder, or condition is pain.

18. The method of claim 17, wherein the compound is administered in combination with morphine.

19. The method of claim 16, wherein the glutamate mediated disease, disorder, or condition is schizophrenia.

20. The method of claim 16, wherein the glutamate mediated disease, disorder, or condition is diabetic neuropathy.

21. A method for effecting a neuronal activity in a mammal, comprising administering to said mammal an effective an effective amount of a compound containing a metabotropic glutamate receptor ligand attached to a metal chelating group.

22. A method for treating a compulsive disorder in a mammal, comprising administering to said mammal an effective an effective amount of a compound containing a metabotropic glutamate receptor ligand attached to a metal chelating group.

23. A method for inhibiting angiogenesis in a mammal, comprising administering to said mammal an effective an

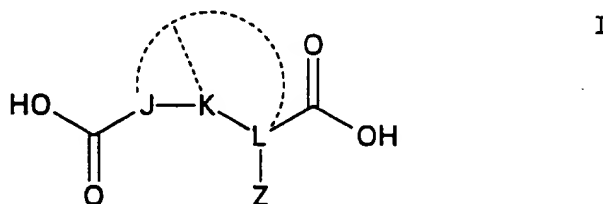


effective amount of a compound containing a metabotropic glutamate receptor ligand attached to a metal chelating group.

24. A method for treating a prostate disease in a mammal, comprising administering to said mammal an effective amount of a compound containing a metabotropic glutamate receptor ligand attached to a metal chelating group.

25. A pharmaceutical composition comprising:

(i) an effective amount of a compound of formula I:



or a pharmaceutically acceptable equivalent, wherein

either J and K are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 3-7 membered saturated or unsaturated heterocyclic or carbocyclic ring,

or J, K, and L are taken together with one or more additional atoms independently selected from the group consisting of C, O, S, and N in chemically reasonable substitution patterns to form a 4-8 membered saturated or unsaturated, mono-, bi-, or tricyclic, hetero- or carbocyclic ring structure;

Z is a metal chelating group; and

(ii) a pharmaceutically acceptable carrier.